## SEARCH REQUEST FORM

Sc	ientific and Tech	inical Information (	Center		•
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Requester's Full Name: //	stromen	Examiner # :_	69630 Da	te: <u>4//0/</u>	<u> </u>
Art Unit: 1651 Phone	Jumber 30 800	732 Serial Nu	mber: 10/0/	17,625	
Requester's Full Name:	1: //30/	Results Format Pref	erred (circle): PA	APER DISK	E-MAIL
If more than one search is subm	itted, please pri	oritize searches in	order of need.		
Please provide a detailed statement of the					
Include the elected species or structures, k	eywords, synonyms,	acronyms, and registry r	numbers, and combi	ine with the cond	ept or
utility of the invention. Define any terms known. Please attach a copy of the cover s			pies or relevant cita	itions, authors, e	ic, 11
Title of Invention:				•	
				_	
Inventors (please provide full names): _		····			
	· ·			<u> </u>	
Earliest Priority Filing Date:	7/3/1996				
*For Sequence Searches Only* Please include appropriate serial number.	le all pertinent informa	ntion (parent, child, divisio	nal, or issued patent	numbers) along w	ith the
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STAFF USE ONLY	Type of Search	Vendor	s and cost where a	pplicable	
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Date Completed:	Litigation				_
Searcher Prep & Review Time:	Fulltext				_
Clerical Prep Time:	Patent Family	WWW/Internet			_

PTO-1590 (8-01)

Online Time: \_

## BioTech-Chem Library Search Results Feedback Form (Optional)

mary.hale@uspto.gov.



The search results generated for your recent request are attached. If you have any questions or comments (compliments or complaints) about the scope or the results of the search, please contact *the BioTech-Chem searcher* who conducted the search *or contact*:

Mary Hale, Supervisor, 308-4258 CM-1 Room 1E01

۶	I am an examiner in Workgroup: (Example: 1610)
٦	Relevant prior art found, search results used as follows:
	102 rejection
	103 rejection
	Cited as being of interest.
	Helped examiner better understand the invention.
	Helped examiner better understand the state of the art in their technology.
	Types of relevant prior art found:
	Foreign Patent(s)
	Non-Patent Literature (journal articles, conference proceedings, new product announcements etc.)
2	Relevant prior art not found:
	Results verified the lack of relevant prior art (helped determine patentability).
	Search results were not useful in determining patentability or understanding the invention.
Oth	er Comments:

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L100 ANSWER 1 OF 15 HCAPLUS COPYRIGHT 2003 ACS

AN 2003:117975 HCAPLUS

DN 138:166253

- TI Media and methods for cultivation and detection of fastidious microorganisms
- IN Breitschwerdt, Edward B.; Sontakke, Sushama
- PA North Carolina State University, USA
- SO PCT Int. Appl., 57 pp.

CODEN: PIXXD2

DT Patent

LA English

IC ICM C12N

CC 9-16 (Biochemical Methods)
Section cross-reference(s): 10, 13, 14

FAN.CNT 1

L'MIA'	> 1 N T	1																
	PATENT NO.			KIND DATE		APPLICATION NO.				ο.	DATE							
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PI WO 2003012058			A2 20030213				WO 2002-US24329 20					20020	0020731					
		W:	ΑE,	AG,	AL,	AM,	AT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,
			CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,
			GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,
			LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	ΜZ,	NO,	NZ,	OM,	PH,
			PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,	TM,	TN,	TR,	TT,	TZ,
			UA,	UG,	US,	UZ,	VN,	YU,	ZA,	ZM,	ZW,	AM,	ΑZ,	BY,	KG,	ΚZ,	MD,	RU,
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			CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	ΙE,	IT,	LU,	MC,	NL,
			PT,	SE,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,
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AB	The	e pre	sent	inv	enti	on p	rovi	des (	cult	ure 1	nedia	a and	d					

AB The present invention provides culture media and methods for culturing organisms, preferably microorganisms, more preferably fastidious microorganisms. Also provided are methods of isolating and detecting organisms using the inventive culture media. Microorganisms were isolated from various clin. samples from canines and felines and grown on D2 insect growth medium.

ST culture media cultivation detection fastidious

microorganism; cat fastidious microorganism clin sample culture; dog fastidious microorganism clin sample culture ΤT Fatty acids, biological studies RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (Me esters, of cod liver oil; media and methods for cultivation and detection of fastidious microorganisms) TT Proteobacteria (alpha group, Rasbo bacterium; media and methods for cultivation and detection of fastidious microorganisms) TT Anemia (disease) (autoimmune hemolytic anemia, fastidious microorganism assocd. with; media and methods for cultivation and detection of fastidious microorganisms) Samples IT (biol., of mammal; media and methods for cultivation and detection of fastidious microorganisms) TΤ Disease, animal (cat scratch disease, fastidious microorganism assocd. with; media and methods for cultivation and detection of fastidious microorganisms) IT Nervous system (central, disease, fastidious microorganism assocd. with; media and methods for cultivation and detection of fastidious microorganisms) IT Chemistry (chem. compds., reducing growth or viability of fastidious microorganism, identification of; media and methods for cultivation and detection of fastidious microorganisms) Fatigue, biological IT (chronic fatigue syndrome, fastidious microorganism assocd. with; media and methods for cultivation and detection of fastidious microorganisms) IT Liver, disease (chronic, fastidious microorganisms culture from; media and methods for cultivation and detection of fastidious microorganisms) IT (chyle, feline chylothorax fluid, fastidious microorganisms culture from cats with; media and methods for cultivation and detection of fastidious microorganisms) IT Mammalia (culture medium contg. lipids of brain tissue of; media and methods for cultivation and detection of fastidious microorganisms) IT (culture medium contg. lipids of mammalian tissue of; media and methods for cultivation and detection of fastidious microorganisms) IT Antioxidants Reducing agents (culture medium contg.; media and methods for cultivation and detection of fastidious microorganisms) IT Amino acids, biological studies Hemins Lipids, biological studies Minerals, biological studies Nucleotides, biological studies Vitamins RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (culture medium contq.; media and methods for cultivation and detection of fastidious microorganisms) TT

Proteins

RL: BSU (Biological study, unclassified); BIOL (Biological study) (culture medium free of; media and methods for cultivation and detection of fastidious microorganisms) Liquids IΤ Solids (culture medium; media and methods for cultivation and detection of fastidious microorganisms) IT Prostate gland Urinary tract (disease, fastidious microorganism assocd. with; media and methods for cultivation and detection of fastidious microorganisms) TΤ Pleura (effusion, fastidious microorganisms culture from dogs; media and methods for cultivation and detection of fastidious microorganisms) IT Body fluid (effusion; media and methods for cultivation and detection of fastidious microorganisms) IT Heart, disease (endocarditis, fastidious microorganisms culture from canines or felines with culture-neg.; media and methods for cultivation and detection of fastidious microorganisms) IT Yeast (ext.; media and methods for cultivation and detection of fastidious microorganisms) TΤ Cystic fibrosis Heart, disease Hypertension Kidney, disease Liver, disease Mastitis Neoplasm (fastidious microorganism assocd. with; media and methods for cultivation and detection of fastidious microorganisms) IT Arachnida Insecta (fastidious microorganism borne by; media and methods for cultivation and detection of fastidious microorganisms) ΙT (fastidious microorganism impaired by; media and methods for cultivation and detection of fastidious microorganisms) IT Canidae Cat (Felis catus) Cattle Dog (Canis familiaris) Goat Horse (Equus caballus) Lagomorpha Primates Rodentia Sheep Swine (fastidious microorganism pathogenic in; media and methods for cultivation and detection of fastidious microorganisms) Biological transport ΙT (fastidious microorganism with defects in nutrient; media and methods for cultivation and detection of fastidious microorganisms) ΙT Nutrients (fastidious microorganism with defects in transport of; media and methods for cultivation and detection of fastidious microorganisms) Metabolism, microbial IT (fastidious microorganism with defects in; media and methods for cultivation and detection of fastidious microorganisms)

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Stress, microbial
IT
        (fastidious microorganism; media and methods for cultivation
        and detection of fastidious microorganisms)
IΤ
     Bacteria (Eubacteria)
     Microorganism
        (fastidious; media and methods for cultivation and detection
        of fastidious microorganisms)
     Cod liver oil
TΤ
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (fatty acid Me esters; media and methods for cultivation and
        detection of fastidious microorganisms)
     Disease, animal
IT
        (feline urol. syndrome, fastidious microorganisms culture
        from cats with; media and methods for cultivation and
        detection of fastidious microorganisms)
TT
     Abdomen
     Thorax
        (fluid of; media and methods for cultivation and detection of
        fastidious microorganisms)
TΤ
     Bioassay
        (for compds. binding to fastidious microorganism; media and
        methods for cultivation and detection of fastidious microorganisms)
     Lactalbumins
TT
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (hydrolyzates; media and methods for cultivation and
        detection of fastidious microorganisms)
TT
    Adrenal cortex, disease
        (hyperadrenocorticism, fastidious microorganism assocd: with;
       media and methods for cultivation and detection of fastidious
       microorganisms)
     Protein hydrolyzates
ΙT
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (lactalbumin hydrolyzates; media and methods for cultivation
        and detection of fastidious microorganisms)
IT
     Pathogen
        (mammalian; media and methods for cultivation and detection
        of fastidious microorganisms)
IT
     Aeromonas hydrophila
     Afipia
     Ascitic fluid
     Bacillus clausii
     Bartonella
     Bartonella clarridgeiae
     Bartonella henselae
     Bartonella vinsonii berkhoffii
     Bartonella weissii
     Blood
     Blood plasma
     Blood products
     Blood serum
     Body fluid
     Bordetella bronchiseptica
     Brucella
     Cerebrospinal fluid
     Corynebacterium
       Culture media
     Escherichia coli
     Exudate
     Hydrogenophaga
     Lymph
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Mycobacterium kansasii Nanobacterium Pleural fluid Proteobacteria Proteus mirabilis Pseudomonas aeruginosa Pseudomonas plecoglossicida Ralstonia pickettii Salmonella typhimurium Sputum Streptococcus pneumoniae Streptococcus thermophilus Synovial fluid Urine (media and methods for cultivation and detection of fastidious microorganisms) Phosphatidylcholines, biological studies RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (media and methods for cultivation and detection of fastidious microorganisms) Encephalitis (meningoencephalitis, granulomatous, culturing synovial fluid of dog with; media and methods for cultivation and detection of fastidious microorganisms) Cell wall (microorganism deficient in; media and methods for cultivation and detection of fastidious microorganisms) (mucus of; media and methods for cultivation and detection of fastidious microorganisms) Arthritis (neutrophilic, culturing synovial fluid of dog with; media and methods for cultivation and detection of fastidious microorganisms) рΗ (of culture medium; media and methods for cultivation and detection of fastidious microorganisms) Diagnosis (of disorder or infection with fastidious microorganism; media and methods for cultivation and detection of fastidious microorganisms) Acids, biological studies RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (org., culture medium contg.; media and methods for cultivation and detection of fastidious microorganisms) Arthritis (polyarthritis, fastidious microorganism assocd. with; media and methods for cultivation and detection of fastidious microorganisms) Filtration (prior to culturing; media and methods for cultivation and detection of fastidious microorganisms) Kidney, disease (protein-losing, fastidious microorganisms culture from; media and methods for cultivation and detection of fastidious microorganisms) Mucus (pulmonary; media and methods for cultivation and detection of fastidious microorganisms) Kidney, disease (pyelonephritis; media and methods for cultivation and detection of fastidious microorganisms) Culture media

ΙT

IT

TΤ

TT

ΙT

ΙT

ΙT

ΙT

IT

TT

ΙT

IT

IT

IT

(selective; media and methods for cultivation and detection of fastidious microorganisms) IT Carbohydrates, biological studies RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (simple, culture medium contg.; media and methods for cultivation and detection of fastidious microorganisms) IT Platelet (blood) (thrombocytopenia, fastidious microorganism assocd. with; media and methods for cultivation and detection of fastidious microorganisms) ITAntibodies RL: ANT (Analyte); BSU (Biological study, unclassified); ANST (Analytical study); BIOL (Biological study) (to fastidious microorganism; media and methods for cultivation and detection of fastidious microorganisms) IT Body fluid (transudate; media and methods for cultivation and detection of fastidious microorganisms) IT Infection (with fastidious microorganism, diagnosis of; media and methods for cultivation and detection of fastidious microorganisms) ΙT Globulins, biological studies RL: ADV (Adverse effect, including toxicity); BSU (Biological study, unclassified); BIOL (Biological study) (.gamma.-, hypergammaglobulinemia, splenomegaly, culturing blood of cat with; media and methods for cultivation and detection of fastidious microorganisms) ΙT 53-59-8, NADP 53-84-9, NAD 1192-20-7D, Homoserine lactone, acyl derivs. RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (culture medium contq.; media and methods for cultivation and detection of fastidious microorganisms) ΙT 50-69-1, Ribose 50-81-7, L-Ascorbic acid, biological studies 50-89-5, Thymidine, biological studies 50-99-7, Dextrose, biological studies 51-35-4, Hydroxy-L-proline **52-90-4**, L-Cysteine, biological studies 53-57-6, .beta.-NADPH 56-40-6, Glycine, biological studies 56-41-7, L-Alanine, biological studies 56-81-5, Glycerol, biological studies 56-84-8, L-Aspartic Acid, biological studies 56-85-9, L-Glutamine, biological studies 56-86-0. L-Glutamic Acid, biological studies 57-48-7, Fructose, biological 57-50-1, Sucrose, biological studies 57-88-5, Cholesterol, biological studies 58-56-0, Pyridoxine hydrochloride 58-85-5 58-95-7, D-.alpha.-Tocopherol Acetate 59-30-3, Folic Acid, biological 59-67-6, Niacin, biological studies 61-90-5, L-Leucine, biological studies 63-68-3, L-Methionine, biological studies 63-91-2, L-Phenylalanine, biological studies 67-03-8, Thiamine hydrochloride 68-19-9, Vitamin B12 67-48-1, Choline Chloride 69-79-4, Maltose 70-18-8, Glutathione, biological studies 70-47-3, 71-00-1, L-Histidine, biological L-Asparagine, biological studies 72-18-4, L-Valine, biological studies 72-19-5, L-Threonine, biological studies 73-22-3, L-Tryptophan, biological studies 73 - 32 - 5, L-Isoleucine, biological studies 77-92-9, Citric Acid, biological 83-88-5, Riboflavin, biological studies 85-61-0, Coenzyme A, biological studies 87-89-8, myo-Inositol 97-67-6, L-Malic Acid

137-08-6, Calcium D-Pantothenate 144-55-8, Sodium Bicarbonate, biological studies 147-85-3, L-Proline, biological studies 150-13-0, PABA 154-87-0, Cocarboxylase 302-84-1, Serine 328-50-7, .alpha.-Ketoglutaric Acid 657-27-2, L-Lysine hydrochloride 838-07-3, 5-Methyl-2'-Deoxycytidine 958-09-8, 2'-Deoxyadenosine 961-07-9, 2'-Deoxyguanosine 1119-34-2, L-Arginine

110-15-6, Succinic Acid, biological studies

127-17-3, Pyruvic Acid,

107-95-9, .beta.-Alanine

biological studies

110-17-8, Fumaric Acid, biological studies

hydrochloride 1310-73-2, **Sodium** hydroxide, biological studies 1336-21-6, Ammonium hydroxide 3992-42-5, 2'-Deoxycytidine 7365-45-9, HEPES monohydrochloride 7447-39-4, Copper chloride 7447-40-7, Potassium (CuCl2), biological studies Chloride, biological studies 7487-88-9, Magnesium Sulfate, biological studies 7512-17-6, N-Acetyl-D-Glucosamine 7558-79-4 7558-80-7, Sodium Phosphate, Monobasic 7646-79-9, Cobalt **chloride**, biological studies 7646-85-7, Zinc 7720-78-7 chloride, biological studies 7786-30-3, Magnesium Chloride, biological studies 9005-65-6, Polyoxyethylenesorbitan Monooleate 9012-36-6, Agarose 10043-52-4, Calcium Chloride, biological studies 10597-89-4, 11104-38-4, Vitamin K1 N-Acetylmuramic Acid 12027-67-7, Ammonium molybdate 30925-07-6, L-Cystine dihydrochloride 69847-45-6, L-Tyrosine 74674-72-9 106392-12-5, Pluronic F-68 disodium salt RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)

(media and methods for cultivation and detection of fastidious microorganisms)

IT 497129-98-3 497129-99-4 497130-00-4 497130-01-5 497130-02-6 497130-03-7 497130-04-8 497130-05-9

RL: PRP (Properties)

(unclaimed sequence; **media** and methods for cultivation and detection of fastidious microorganisms)

IT 50-89-5, Thymidine, biological studies 52-90-4, L-

Cysteine, biological studies

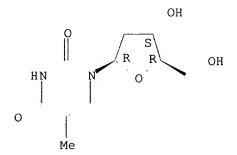
RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)

 $(\mbox{media}\mbox{ and methods for cultivation and detection of }\mbox{fastidious microorganisms})$ 

RN 50-89-5 HCAPLUS

CN Thymidine (8CI, 9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 52-90-4 HCAPLUS

CN L-Cysteine (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L100 ANSWER 2 OF 15 HCAPLUS COPYRIGHT 2003 ACS

AN 2001:353374 HCAPLUS

DN 135:149238

```
TΙ
     Pantothenic acid protects Jurkat cells against
     ultraviolet light-induced apoptosis
     Slyshenkov, V. S.; Piwocka, K.; Sikora, E.; Wojtczak, L.
ΑU
     Nencki Institute of Experimental Biology, Polish Academy of Sciences,
CS
     Warsaw, Pol.
     Free Radical Biology & Medicine (2001), 30(11), 1303-1310
SO
     CODEN: FRBMEH; ISSN: 0891-5849
     Elsevier Science Inc.
PB
DΤ
     Journal
     English
LA
     8-9 (Radiation Biochemistry)
CC
     Human leukemic T lymphocytes (Jurkat cells) were induced to
AΒ
     undergo apoptosis by brief irradn. with UV C light (254 nm).
                                                                     This was
     accompanied by accumulation of lipid peroxidn. products in the form of
     conjugated dienes, a decrease of total glutathione content, and a shift of
     its redox state towards the oxidized form. Preincubation of the cells
     with 1 mM pantothenate resulted in a significant elevation of
     total glutathione content of the cells, reaching its max. level, 160% of
     the control, after 3 h. Similar increase was obsd. after preincubation
     with 5 mM N-acetylcysteine, a known precursor of
     glutathione. Both pantothenic acid and N-
     acetylcysteine alleviated the UV-induced decrease of glutathione
     content, diminished lipid peroxidn., and partly protected the cells
     against apoptosis produced by UV irradn.
ST
     pantothenate UVC apoptosis lipid peroxidn glutathione
     Peroxidation
IΤ
        (lipid; pantothenic acid protection against
        UV-induced apoptosis)
IT
     Antioxidants
     Apoptosis
     Oxidative stress, biological
     Radioprotectants
     UV C radiation
        (pantothenic acid protection against UV-induced
        apoptosis)
ΙT
     Lipids, biological studies
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (peroxidn.; pantothenic acid protection against
        UV-induced apoptosis)
IT
     616-91-1, N-Acetylcysteine
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (pantothenic acid protection against UV-induced
        apoptosis)
ΤТ
     79-83-4, Pantothenic acid
                                  867-81-2,
     Sodium Pantothenate
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (pantothenic acid protection against UV-induced
        apoptosis)
TT
     70-18-8, Glutathione, biological studies
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (pantothenic acid protection against UV-induced
        apoptosis)
              THERE ARE 53 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
(1) Aprahamian, M; Am J Clin Nutr 1985, V41, P578 HCAPLUS
(2) Artom, C; Proc Soc Exp Biol Med 1954, V86, P162 HCAPLUS (3) Beaver, J; Eur J Cell Biol 1995, V68, P47 HCAPLUS
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(4) Borets, V; Vopr Pitan 1983, 1, P45 MEDLINE

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RL: BAC (Biological activity or effector, except adverse); BSU (Biological

(pantothenic acid protection against UV-induced apoptosis)

study, unclassified); BIOL (Biological study)

616-91-1 HCAPLUS RN

IT

L-Cysteine, N-acetyl- (9CI) (CA INDEX NAME)

616-91-1, N-Acetylcysteine

Absolute stereochemistry.

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HS
       NHAC
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        (pantothenic acid protection against UV-induced
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     79-83-4 HCAPLUS
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L100 ANSWER 3 OF 15 HCAPLUS COPYRIGHT 2003 ACS
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     Animal cell culture media containing plant-derived
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IN
     Price, Paul J.; Gorfien, Steve; Danner, Douglas
     Life Technologies, Inc., USA; Price, Paul J.; Gorfien, Steve; Danner,
PA
     Douglas
SO
     PCT Int. Appl., 58 pp.
     CODEN: PIXXD2
DT
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LA
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     ICS A61K035-72; A61K035-78
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     9-11 (Biochemical Methods)
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EP 954563

US 6103529

PRAI US 1996-28197P

JP 2001501830

A1

IE, SI, LT, LV, FI, RO

Α

T2

Ρ

19991110

20000815

20010213

19961010

EP 1997-910043

US 1997-949142

JP 1998-517748

AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,

19971010

19971010

19971010

WO 1997-US18255 W 19971010 AΒ The present invention provides serum-free cell culture media formulations which are capable of supporting the in vitro cultivation of animal cells. The media comprise at least one nutrient of plant derivation, such as at least one plant peptide and/or at least one plant lipid and/or at least one plant fatty acid. The media may further optionally comprise an enzymic digest or ext. of yeast cells. The present invention also provides methods of cultivating animal cells in vitro using these cell culture media formulations. The application of plant-derived nutrients can reduce the use of serum or animal exts. in tissue culture applications. ST animal tissue culture plant derived nutrient IΤ Animal cell line (293; animal cell culture media contg. plant-derived nutrients) ΙT Animal cell line (BHK-21; animal cell culture media contg. plant-derived nutrients) IT · Animal cell line (BHK; animal cell culture media contg. plant-derived nutrients) TΤ Animal cell line (CHO; animal cell culture media contg. plant-derived nutrients) TT Animal cell line (COS-7; animal cell culture media contg. plant-derived nutrients) Animal cell line IT (COS; animal cell culture media contq. plant-derived nutrients) IT Animal cell line (K562; animal cell culture media contg. plant-derived nutrients) IT Animal cell line (M1; animal cell culture media contg. plant-derived nutrients) ΙT Animal cell line (MDBK; animal cell culture media contg. plant-derived nutrients) ΙT Animal cell line (MDCK; animal cell culture media contg. plant-derived nutrients) IT Animal cell line (MRC-5; animal cell culture media contg. plant-derived nutrients) IT Animal cell line (Molt 4; animal cell culture media contg. plant-derived nutrients) ΙT Animal cell line (NS-1; animal cell culture media contg. plant-derived nutrients) IT Animal cell line (PER-C6; animal cell culture media contg. plant-derived nutrients) ITAnimal cell line (Sp2/0; animal cell culture media contg. plant-derived nutrients) IT Animal cell line (Vero; animal cell culture media contg. plant-derived nutrients) IT Animal cell line

(WEHI; animal cell culture media contg.

```
plant-derived nutrients)
    Animal cell line
TT
        (WI-38; animal cell culture media contg.
        plant-derived nutrients)
TΤ
     Animal cell
    Animal tissue culture
   . Bakers' yeast
     Bird (Aves)
       Culture media
     Fish
     HeLa cell
     Hybridoma
     Insect (Insecta)
     Mammal (Mammalia)
     Plant (Embryophyta)
     Potato (Solanum tuberosum)
     Rice (Oryza sativa)
     Soybean (Glycine max)
     Spodoptera
     Trichoplusia
     Virus
     Wheat
     Yeast
        (animal cell culture media contg. plant-derived
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    Amino acids, biological studies
    Mineral elements, biological studies
     Proteins, general, biological studies
     Salts, biological studies
     Trace element nutrients
    Vitamins
     RL: BSU (Biological study, unclassified); BUU (Biological use,
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        (animal cell culture media contg. plant-derived
        nutrients)
ΙT
     Fatty acids, biological studies
     Lipids, biological studies
     Peptides, biological studies
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
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        (animal cell culture media contg. plant-derived
        nutrients)
TΨ
    Nutrients
        (macronutrients; animal cell culture media contg.
        plant-derived nutrients)
IT
     Nutrients
        (micronutrients; animal cell culture media contg.
        plant-derived nutrients)
IT
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     RL: BAC (Biological activity or effector, except adverse); BPR (Biological
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        (animal cell culture media contg. plant-derived
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     L-Aspartic acid, biological studies 56-86-0, L-Glutamic acid, biological
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59-30-3, Folic acid, biological studies 59-43-8, Thiamine, biological 59-67-6, Nicotinic acid, biological studies 60-18-4, L-Tyrosine, biological studies 60-33-3, Linoleic acid, biological 61-90-5, L-Leucine, biological studies 63-68-3, L-Methionine, studies biological studies 63-91-2, L-Phenylalanine, biological studies 65-23-6, Pyridoxine 66-22-8, Uracil, biological studies Pyridoxal 67-48-1, Choline chloride 68-19-9, 68-94-0, Hypoxanthine 69-89-6, Xanthine 70-18-8, Vitamin B12 Glutathione, biological studies 70-47-3, L-Asparagine, biological 71-00-1, L-Histidine, biological studies 72-18-4, L-Valine, studies 72-19-5, L-Threonine, biological studies biological studies 73-22-3, L-Tryptophan, biological studies 73-32-5, L-Isoleucine, biological 83-88-5, Riboflavin, 74-79-3, L-Arginine, biological studies biological studies 87-89-8, myo-Inositol 98-92-0, Niacinamide 112-80-1, Oleic acid, biological studies 110-60-1, Putrescine 112-85-6, Behenic acid 112-86-7, Erucic acid 113-24-6, Sodium pyruvate 124-07-2, Caprylic acid, biological studies 127-47-9, Vitamin 137-08-6 141-43-5, Ethanolamine, biological studies A acetate 143-07-7, Lauric acid, biological studies 143-74-8, Phenol red 144-55-8, Sodium bicarbonate, biological studies 147-85-3, L-Proline, biological studies 150-13-0 321-30-2, Adenine sulfate 334-48-5, Capric acid 373-49-9, Palmitoleic acid 463-40-1, Linolenic 506-30-9, Arachidic acid 533-67-5, Deoxyribose 543-80-6, Barium 544-63-8, Myristic acid, biological studies 557-59-5, acetate Lignoceric acid 1071-23-4, Phosphoethanolamine 1077-28-7, 1,2-Dithiolane-3-pentanoic acid 1310-53-8, Germanium oxide, biological 2002-24-6, Ethanolamine hydrochloride 6834-92-0, 7365-45-9, HEPES 7429-90-5, Sodium silicate (Na2SiO3) Aluminum, biological studies 7439-89-6, Iron, biological studies 7439-96-5, Manganese, biological studies 7439-98-7, Molybdenum, biological studies 7440-02-0, Nickel, biological studies 7440-17-7, Rubidium, biological studies 7440-21-3, Silicon, biological 7440-31-5, Tin, 7440-22-4, Silver, biological studies 7440-39-3, biological studies 7440-32-6, Titanium, biological studies Barium, biological studies 7440-43-9, Cadmium, biological studies 7440-47-3, Chromium, biological studies 7440-48-4, Cobalt, biological 7440-50-8, Copper, biological studies 7440-56-4, Germanium, biological studies 7440-62-2, Vanadium, biological studies 7440-66-6, Zinc, biological studies 7440-67-7, Zirconium, biological studies 7447-40-7, Potassium chloride, biological studies 7487-88-9, Magnesium sulfate, biological studies 7550-45-0, Titanium chloride, biological studies 7553-56-2, Iodine, 7558-79-4, Disodium hydrogen phosphate biological studies 7558-80-7, Sodium dihydrogen phosphate 7646-79-9, 7647-14-5, Sodium Cobalt chloride, biological studies chloride, biological studies 7681-11-0, Potassium iodide, biological studies 7681-49-4, Sodium fluoride, biological studies 7699-43-6, Zirconium dichloride monoxide 7720-78-7, 7726-95-6, Bromine, biological studies Iron sulfate (FeSO4 7733-02-0, Zinc sulfate 7758-02-3, Potassium bromide, biological studies 7758-98-7, Copper sulfate, biological studies 7772-99-8, Tin 7761-88-8, Silver nitrate, biological studies chloride, biological studies 7773-01-5, Manganese 7782-41-4, Fluorine, biological studies chloride 7782-49-2, 7783-00-8, Selenious acid 7786-30-3, Selenium, biological studies 7786-81-4, Magnesium chloride, biological studies 7791-11-9, Rubidium chloride, biological Nickel sulfate 9004-10-8, Insulin, biological studies 10043-01-3, Aluminum studies sulfate 10043-52-4, Calcium chloride, biological 10124-36-4, Cadmium sulfate 10421-48-4, Ferric nitrate studies 12027-67-7, Ammonium molybdate ((NH4)6Mo7024) 13718-26-8, Sodium 14489-25-9, Chromium sulfate 15431-40-0, vanadate (NaVO3) Magnesium ascorbate 28633-45-6, Ferric citrate

RL: BSU (Biological study, unclassified); BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (animal cell culture media contg. plant-derived nutrients)

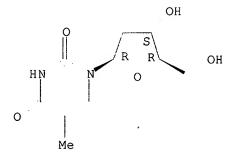
RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD RE

- (1) Anon; GIBCOBRL LIFE TECHNOLOGIES CATALOGUE AND REFERENCE GUIDE 1993, P1
- (2) Durnford; US 5324524 A 1994 HCAPLUS
- (3) Maurer, H; Animal Cell Culture, A Practical Approach 1986, P13
- (4) Sakata; US 5318906 A 1994 HCAPLUS
- (5) Soken; JP 07165524 A 1995
- (6) Wyss, C; Somatic Cell Genetics 1979, V5(1), P23 HCAPLUS
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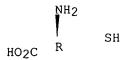
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Absolute stereochemistry.



RN 52-90-4 HCAPLUS CN L-Cysteine (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 7439-89-6 HCAPLUS CN Iron (7CI, 8CI, 9CI) (CA INDEX NAME)

Fe

L100 ANSWER 4 OF 15 HCAPLUS COPYRIGHT 2003 ACS

AN 1998:176040 HCAPLUS

DN 128:228255

TI Assessment of intracellular **cysteine** and glutathione concentrations

IN Crawford, J. Fred

PA Research Development Foundation, USA

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SO
     PCT Int. Appl., 36 pp.
     CODEN: PIXXD2
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     English
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     ICM C120001-02
     ICS C12N005-00
CC
     9-11 (Biochemical Methods)
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    A medium and method for culturing lymphocytes are provided for
AB
    detq. intracellular concn. of glutathione or cysteine in human
     lymphocytes to provide biochem. anal. of an individual's
     capability of dealing with oxidative stress. The medium is a buffered
     serum-free soln. having a pH of from about 6.8 to 7.6
     and contg. a carbohydrate which is glucose or a compd. capable of
     producing glucose in lymphocytes, pantothenic
     acid, choline or a substance capable of producing
     choline in lymphocytes, inorg. ions including
     chloride, phosphate, calcium,
    magnesium, potassium, sodium and iron
     , L-Buthionine-[S.R.]-Sulfoximine, deionized water and a mitogen to
     stimulate lymphocytes. When detg. cysteine concn.,
     the medium addnl. contains N-Acetyl-L Cysteine
     and Cumene Hydroperoxide. The method is carried out
     by inoculating the culture medium with lymphocytes from an
     individual, incubating the lymphocytes in the medium and
     comparing the response of the lymphocytes with an av. response
     of lymphocytes from a control group of individuals.
ST
     intracellular cysteine glutathione concn
    Animal tissue culture
ΙT
     Mitogens
     Oxidative stress, biological
        (assessment of intracellular cysteine and glutathione
        concns.)
IT
     Amino acids, biological studies
     Antioxidants
     Carbohydrates, biological studies
     Vitamins
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (assessment of intracellular cysteine and glutathione
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        concns.)
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             56-45-1, L-Serine, biological studies 58-05-9, Folinic acid
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                      59-30-3, Folic acid, biological studies
     58-85-5, Biotin
                                                                 59-43-8,
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     acid, biological studies 60-18-4, Tyrosine, biological studies
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    Adenine, biological studies 79-83-4, Pantothenic
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                        443-79-8, Isoleucine 616-91-1, N-
                         4998-57-6, Histidine
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     studies 7440-09-7, Potassium, biological studies
    7440-23-5, Sodium, biological studies 7440-70-2
     , Calcium, biological studies
                                     7732-18-5, Water, biological
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        (assessment of intracellular cysteine and glutathione
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RE.CNT
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(1) Bounous; US 5290571 A 1994 HCAPLUS
(2) Darfler; US 4927762 A 1990 HCAPLUS
(3) Griffith; US 5171885 A 1992 HCAPLUS
(4) Ponting; US 5405772 A 1995 HCAPLUS
(5) Torishima; US 5326699 A 1994
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HS-CH_2-CH-CO_2H
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     62-49-7, Choline 79-83-4, Pantothenic
    acid 80-15-9, Cumene Hydroperoxide
     616-91-1, N-Acetyl-L-Cysteine
    7439-89-6; Iron, biological studies 7439-95-4,
    Magnesium, biological studies 7440-09-7,
    Potassium, biological studies 7440-23-5, Sodium
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    RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
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(assessment of intracellular cysteine and glutathione

concns.)

RN 62-49-7 HCAPLUS

CN Ethanaminium, 2-hydroxy-N, N, N-trimethyl- (9CI) (CA INDEX NAME)

 $Me3^{+}N^{-}CH2^{-}CH2^{-}OH$ 

RN 79-83-4 HCAPLUS

CN .beta.-Alanine, N-[(2R)-2,4-dihydroxy-3,3-dimethyl-1-oxobutyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).

OH

 $^{\rm H}_{
m N}$   $^{\rm R}$  OH  $^{\rm OH}$ 

RN 80-15-9 HCAPLUS

CN Hydroperoxide, 1-methyl-1-phenylethyl (9CI) (CA INDEX NAME)

O- OH | Me- C- Me | Ph

RN 616-91-1 HCAPLUS CN L-Cysteine, N-acetyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

HS' R CO2H

RN 7439-89-6 HCAPLUS

CN Iron (7CI, 8CI, 9CI) (CA INDEX NAME)

Fe

RN 7439-95-4 HCAPLUS

CN Magnesium (8CI, 9CI) (CA INDEX NAME)

Mg

RN 7440-09-7 HCAPLUS

CN Potassium (8CI, 9CI) (CA INDEX NAME)

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7440-23-5 HCAPLUS
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      7440-70-2 HCAPLUS
RN
CN
      Calcium (8CI, 9CI) (CA INDEX NAME)
Ca
      14265-44-2 HCAPLUS
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      Phosphate (8CI, 9CI)
                               (CA INDEX NAME)
CN
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-o- p- o-
    0-
      16887-00-6 HCAPLUS
RN
      Chloride (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
CN
C1-
L100 ANSWER 5 OF 15 HCAPLUS COPYRIGHT 2003 ACS
      1998:163670 HCAPLUS
AN
DN
      128:215266
      Serum-free mammalian cell culture medium, and uses
TΙ
      thereof
      Gorfien, Stephen F.; Fike, Richard M.; Dzimian, Joyce L.; Godwin, Glenn
IN
      P.; Price, Paul J.; Epstein, David A.; Gruber, Dale; McClure, Don
      Life Technologies, Inc., USA; Gorfien, Stephen F.; Fike, Richard M.;
PA
      Dzimian, Joyce L.; Godwin, Glenn P.; Price, Paul J.; Epstein, David A.;
      Gruber, Dale; McClure, Don
SO
      PCT Int. Appl., 121 pp.
      CODEN: PIXXD2
DΤ
      Patent
LA
      English
      ICM C12N005-00
ICS C12N005-02; C12N005-06; C12N005-08; C12N005-10
CC
      9-11 (Biochemical Methods)
FAN.CNT 1
      PATENT NO.
                          KIND
                                  DATE
                                                    APPLICATION NO.
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                                 19980305
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               PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
           RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA,
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GN, ML, MR, NE, SN, TD, TG

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AU 9743305
                       A1
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                                            EP 1997-941382
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            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
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                            20001226
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PRAI US 1996-22881P
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     US 1997-56829P
                            19970822
                       Α
     US 1997-920875
                            19970829
     WO 1997-US15296
                       W
                            19970902
     The present invention provides a cell culture medium formulation that
AΒ
     supports the in vitro cultivation, particularly in suspension, of
    mammalian cells, particularly epithelial cells and fibroblast cells, and
    methods for cultivating mammalian cells in suspension in vitro, using
     these media. The media comprise a basal medium and a polyanionic or
     polyanionic compd., preferably a polysulfonated or polysulfated compd.,
     and more preferably dextran sulfate. The present invention also provides
     chem. defined, protein-free eukaryotic cell culture media comprising an
     iron chelate and zinc, which is capable of supporting the growth
     (and particularly the high-d. growth of mammalian cells) in suspension
     culture, increasing the level of expression of recombinant protein in
     cultured cells, and/or increasing virus prodn. in cultured cells.
     serum mammal cell culture medium
ST
ΙT
    Uterus
    Uterus
        (cervix, epithelium; serum-free mammalian cell
        culture medium, and uses thereof)
IT
        (chem. compds., Polyanionic; serum-free mammalian
        cell culture medium, and uses thereof)
IT
     Chemistry
        (chem. compds., Polysulfated; serum-free mammalian
        cell culture medium, and uses thereof)
ΙT
        (chem. compds., Polysulfonated; serum-free
       mammalian cell culture medium, and uses thereof)
ΙT
     Bronchi
     Kidney
    Trachea (anatomical)
        (epithelium; serum-free mammalian cell culture
       medium, and uses thereof)
ΙT
        (keratinocyte; serum-free mammalian cell culture
       medium, and uses thereof)
    Animal cell
IT
        (mammalian; serum-free mammalian cell culture
       medium, and uses thereof)
ΙT
        (retina, epithelium; serum-free mammalian cell
        culture medium, and uses thereof)
IT
    Animal
    Animal tissue culture
    Blood serum
    Buffers
    Culture media
     Embryo, animal
     Epithelium
     Fibroblast
     Kidney
     Plant (Embryophyta)
     Rice (Oryza sativa)
     Soybean (Glycine max)
     Virus
```

Yeast

(serum-free mammalian cell culture medium, and uses thereof) ΙT Amino acids, biological studies Carbohydrates, biological studies Cytokines Lipids, biological studies Peptides, biological studies Proteoglycans, biological studies Salts, biological studies Transferrins Vitamins RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (serum-free mammalian cell culture medium, and uses thereof) ΤT Transferrins RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (substitute; serum-free mammalian cell culture medium, and uses thereof) IT 50-99-7, Glucose, biological studies 52-90-4, Cysteine , biological studies 56-40-6, Glycine, biological studies 56-41-7L-Alanine, biological studies 56-45-1, Serine, biological studies 56-84-8, Aspartic acid, biological studies 56-85-9, Glutamine, 56-86-0, Glutamic acid, biological studies biological studies 56-87-1, Lysine, biological studies 58-85-5, Biotin 59-30-3, Folic acid, biological studies 59-43-8, Thiamine, biological studies 60-18-4, Tyrosine, biological studies 60-33-3, Linoleic acid, biological studies 61-90-5, Leucine, biological studies 63-68-3, Methionine, biological studies 63-91-2, Phenylalanine, biological studies 65-23-6, Pyridoxine 67-48-1, Choline chloride 68-19-9, Vitamin b12 70-47-3, Asparagine, biological studies 71-00-1, Histidine, biological studies 72-18-4, Valine, biological studies 72-19-5, Threonine, biological studies 73-22-3, Tryptophan, biological studies 73-32-5, 74-79-3, Arginine, biological studies Isoleucine, biological studies 83-88-5, Riboflavin, biological studies 87-89-8, i-Inositol Niacinamide 110-60-1, Putrescine 113-24-6, Sodium pyruvate 137-08-6, Calcium pantothenate 141-43-5, 143-74-8, Phenol red 144-55-8, Ethanolamine, biological studies Carbonic acid monosodium salt, biological studies 147-85-3, Proline, 7365-45-9, Hepes **7439-89-6D**, Iron biological studies , chelates, biological studies 7439-95-4D, Magnesium, salts, biological studies 7439-96-5D, Manganese, salts, biological studies 7440-62-2D, Vanadium, salts, biological studies 7440-66-6, 7440-66-6D, Zinc, salts, biological studies Zinc, biological studies 7440-70-2D, Calcium, salts, biological studies 7447-40-7, Potassium chloride (KCl), biological studies 7558-79-4 7647-14-5, Sodium chloride, 7782-49-2D, Selenium, salts, biological studies biological studies 9004-10-8D, Insulin, substitute, 9004-10-8, Insulin, biological studies 9005-49-6, Heparin, biological studies biological studies 9007-28-7, 9050-30-0, Heparan Chondroitin sulfate 9042-14-2, Dextran sulfate 10421-48-4 24967-94-0, Dermatan sulfate 57828-26-9, Lipoic sulfate 106392-12-5, Pluronic f68 140207-93-8, Pentosan sulfate RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (serum-free mammalian cell culture medium, and uses thereof) RE.CNT THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD

(1) Bertheussen; US 5045454 A 1991 HCAPLUS

(2) Inlow; US 5024947 A 1991

RF.

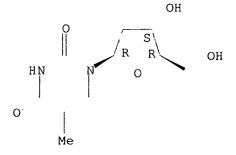
(3) Maurer, H; Animal cell culture:a practical approach 1986, P13

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(4) Mignot; US 5422250 A 1995 HCAPLUS
(5) Skelnik; US 4959319 A 1990
(6) Smithkline Beecham Corporation; WO 9205246 Al 1992 HCAPLUS
(7) Stockinger; US 5063157 A 1991 HCAPLUS
    52-90-4, Cysteine, biological studies 7439-89-6D
     , Iron, chelates, biological studies 7439-95-4D,
    Magnesium, salts, biological studies 7440-70-2D,
    Calcium, salts, biological studies
    RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (serum-free mammalian cell culture medium, and uses
        thereof)
RN
     52-90-4 HCAPLUS
CN
    L-Cysteine (9CI) (CA INDEX NAME)
Absolute stereochemistry.
      NH2
            SH
     R
HO<sub>2</sub>C
    7439-89-6 HCAPLUS
RN
    Iron (7CI, 8CI, 9CI) (CA INDEX NAME)
CN
Fe
    7439-95-4 HCAPLUS
RN
    Magnesium (8CI, 9CI) (CA INDEX NAME)
CN
Mg
RN
    7440-70-2 HCAPLUS
    Calcium (8CI, 9CI) (CA INDEX NAME)
CN
Ca
L100 ANSWER 6 OF 15 HCAPLUS COPYRIGHT 2003 ACS
    1998:105996 HCAPLUS
ΑN
     128:138343
DN
     Chondrocyte media formulations and culture procedures
TΙ
     McPherson, John M.; Yaeger, Peter C.; Brown, Marie E.; Hanlon, James G.;
IN
     Binette, Francois
     Genzyme Corporation, USA; McPherson, John M.; Yaeger, Peter C.; Brown,
PA
     Marie E.; Hanlon, James G.; Binette, Francois
SO
     PCT Int. Appl., 59 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     English
     ICM C12N005-00
IC
     ICS C12N005-06
CC
     9-11 (Biochemical Methods)
FAN.CNT 1
                                           APPLICATION NO.
     PATENT NO.
                      KIND DATE
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         RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
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                       Ρ
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                       Ρ
     US 1996-22711P
                            19960726
    WO 1997-US13140
                      W
                            19970725
    One object of the present invention is based upon the development and use
AB
     of a serum-free defined cell culture
    medium comprising a supplement mixt., a component mixt., a vitamin
    mixt., an inorg. salt mixt. and amino acid mixt. that avoids the problems
     inherent in the use of serum. In particular, the defined medium
     is useful in culturing fibroblasts, esp. chondrocytes. Another
     object of the present invention is to claim a method of enhancing the
    differentiation of chondrocytes and enhancing the synthesis of a cartilage
     specific matrix using tumor growth factor beta (TGF-.beta.). Another
     object of the present invention is to claim a method of enhancing the
     differentiation of chondrocytes using the combination of TGF-.beta. and
ST
     chondrocyte media formulation culture
IT
    Animal tissue culture
     Blood serum
     Cell differentiation
     Chondrocyte
     Fibroblast
        (chondrocyte media formulations and culture
       procedures)
IT
     Amino acids, biological studies
     Cartilage
     Fibronectins
     Salts, biological studies
     Vitamins
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (chondrocyte media formulations and culture
       procedures)
TT
     Transforming growth factors
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (.beta.-; chondrocyte media formulations and culture
       procedures)
IT
     50-23-7, Hydrocortisone 50-89-5, Thymidine, biological studies
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609-36-9, Proline 617-45-8, Aspartic acid 617-65-2, Glutamic acid 3130-87-8, Asparagine **3374-22-9**, **Cysteine** 4998-57-6, 7200-25-1, Arginine 6899-04-3, Glutamine 7447-40-7, Histidine Potassium chloride (KCl), biological studies 7487-88-9, Sulfuric acid magnesium salt (1:1), biological 7558-79-4 7558-80-7 7647-14-5, **Sodium** 7720-78-7 7758-98-7, 7733-02-0 chloride, biological studies Sulfuric acid copper(2+) salt (1:1), biological studies 7786-30-3, Magnesium chloride (MgCl2), biological studies 10043-52-4, Calcium chloride (CaCl2), biological 10124-37-5 10421-48-4 13463-67-7, Its, biological studies 61912-98-9, IGF 62031-54-3, Fgf 57828-26-9, Lipoic acid RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (chondrocyte media formulations and culture procedures) IT 50-89-5, Thymidine, biological studies 3374-22-9, Cysteine RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (chondrocyte media formulations and culture procedures) RN 50-89-5 HCAPLUS CN Thymidine (8CI, 9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 3374-22-9 HCAPLUS CN Cysteine (9CI) (CA INDEX NAME)

NH2 | HS-CH2-CH-CO2H

L100 ANSWER 7 OF 15 HCAPLUS COPYRIGHT 2003 ACS ΑN 1998:31414 HCAPLUS DN 128:72646 Biochemical analysis of antioxidant function TI IN Crawford, J. Fred; Bucci, Luke PA Research Development Foundation, USA PCT Int. Appl., 32 pp. SO CODEN: PIXXD2 DT Patent English LΑ ICM C12Q001-08 TC ICS C12N005-00; C12N005-02; C12N001-38 CC 9-11 (Biochemical Methods)

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                      KIND DATE
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                                                            DATE
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                            19971224
                                           WO 1997-US10328 19970618
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PRAI US 1996-665941
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     WO 1997-US10328
                      W
                            19970618
     The present invention provides a cell culture medium useful for a biochem.
AB
     anal. of antioxidant function in human lymphocytes, said medium
     comprising, a buffered, serum-free soln. contg. the
     following ingredients: a carbohydrate selected from the group consisting
     of glucose and a compd. biol. capable of producing glucose in the cells; a
     biol. usable form of pantothenic acid; choline
     or a biol. usable form of a substance capable of producing choline
     in the cells; inorg. ions comprising chloride, phosphate
     , calcium, magnesium, potassium,
     sodium; and iron in a biol. utilizable form,
     cumene hydroperoxide, deionized water, and a mitogen in
     an amt. effective to stimulate the lymphocytes being assayed; said
     buffered, serum-free soln. having a pH from about 6.8
     to 7.6, said cell culture medium characterized by being effective to det.
     nutritional deficiencies, inadequacies, and imbalances and to biochem.
     analyze antioxidant function of the lymphocytes. Also provided is a
     method of biochem. analyzing cellular antioxidant function in an
     individual comprising the steps of: inoculating the cell culture medium of
     the present invention with lymphocytes from said individual; incubating
     the inoculated cell culture medium; and comparing the response of the
     lymphocytes with an av. response of lymphocytes from a control group of
     individuals.
ST
     biochem analysis antioxidant function
IT
     Animal tissue culture
     Antioxidants
     Blood serum
     Culture media
     Ions
     Lymphocyte
     Mitogens
        (biochem. anal. of antioxidant function)
TI
     Buffers
     Nutrition, animal
     RL: ANT (Analyte); ANST (Analytical study)
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(biochem. anal. of antioxidant function)

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ΙT
    Amino acids, biological studies
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (biochem. anal. of antioxidant function)
ΙT
     Carbohydrates, biological studies
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (biochem. anal. of antioxidant function)
ΙT
    Vitamins
    RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (biochem. anal. of antioxidant function)
ΙT
    Analysis
        (biochem.; biochem. anal. of antioxidant function)
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    Cysteine, biological studies
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    Nicotinamide
    Iron, biological studies 7439-95-4, Magnesium,
    biological studies 7440-09-7, Potassium, biological
    studies 7440-23-5, Sodium, biological studies
    7440-70-2, Calcium, biological studies
                                              7732-18-5,
    Water, biological studies
                                8059-24-3, Vitaminb6 14265-44-2,
    Phosphate, biological studies 16887-00-6,
    Chloride, biological studies
    RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (biochem. anal. of antioxidant function)
ΤТ
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    80-15-9, Cumene hydroperoxide
    7439-89-6, Iron, biological studies 7439-95-4,
    Magnesium, biological studies 7440-09-7,
    Potassium, biological studies 7440-23-5, Sodium
     , biological studies 7440-70-2, Calcium, biological
    studies 14265-44-2, Phosphate, biological studies
    16887-00-6, Chloride, biological studies
    RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (biochem. anal. of antioxidant function)
RN
    52-90-4 HCAPLUS
    L-Cysteine (9CI)
                      (CA INDEX NAME)
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Absolute stereochemistry.

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NH2
HO2C R
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RN 62-49-7 HCAPLUS

CN Ethanaminium, 2-hydroxy-N,N,N-trimethyl- (9CI) (CA INDEX NAME)

 ${
m Me3}^{+}{
m N}-{
m CH}_{2}-{
m CH}_{2}-{
m OH}$ 

RN 79-83-4 HCAPLUS

CN .beta.-Alanine, N-[(2R)-2,4-dihydroxy-3,3-dimethyl-1-oxobutyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).

$$^{\rm OH}_{\rm N}$$
  $^{\rm R}_{\rm OH}$   $^{\rm OH}_{\rm OM}$ 

RN 80-15-9 HCAPLUS

CN Hydroperoxide, 1-methyl-1-phenylethyl (9CI) (CA INDEX NAME)

RN 7439-89-6 HCAPLUS

CN Iron (7CI, 8CI, 9CI) (CA INDEX NAME)

Fe

RN 7439-95-4 HCAPLUS

CN Magnesium (8CI, 9CI) (CA INDEX NAME)

Mg

RN 7440-09-7 HCAPLUS

CN Potassium (8CI, 9CI) (CA INDEX NAME)

K

RN 7440-23-5 HCAPLUS

CN Sodium (8CI, 9CI) (CA INDEX NAME)

Na

RN 7440-70-2 HCAPLUS

CN Calcium (8CI, 9CI) (CA INDEX NAME)

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Ca
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14265-44-2 HCAPLUS RN Phosphate (8CI, 9CI) (CA INDEX NAME) CN

RN 16887-00-6 HCAPLUS

CN Chloride (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

C1-

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L100 ANSWER 8 OF 15 HCAPLUS COPYRIGHT 2003 ACS
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1997:761704 HCAPLUS AN

DN 128:45595

TIMethod for serum-free culture of human

vascular endothelial cells

Katsuen, Susumu; Ohshima, Kunihiro; Yamamoto, Ryohei; Nishino, Toyokazu Kurashiki Boseki Kabushiki Kaisha, Japan IN

PA

U.S., 11 pp. CODEN: USXXAM

DT Patent

LA English

ICM C12N005-00

ICS C12N011-00; C12N011-02; C12N011-08

9-11 (Biochemical Methods)

FAN.CNT 1

					DATE	
	PATENT NO.	KIND	DATE	APPLICATION NO.		
ΡI	US 5691203	A	19971125	US 1993-128225	19930929	
	JP 07008273	A2	19950113	JP 1993-141984	19930614	
PRAI	JP 1993-141984		19930614			
AB	Animal adhesive	cells,	particularly	human vascular end	othelial c	
	cultured in seru	m-free	condition by	coating at		

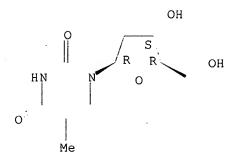
cells, are cultured in serum-free condition by coating at least one polymer having cell adhesive activity on an inner surface of a culture vessel or surface of a carrier for cell culture, and culturing the cells in the coated vessel or with the coated carrier using a serum-free medium for animal cell culture contq. isolated serum albumin, and preferably also transferrin. The polymer is a synthetic polymer modified with a peptide having cell adhesive activity or a natural polymer having cell adhesive activity or a combination thereof. Preferably, the peptide is RGDV, RGDS, RGDN, DGEA or YIGSR and the natural polymer is collagen, gelatin, keratin, fibronectin, vitronectin or laminin. A preferred medium for culturing human vascular endothelial cells is basal medium MCDB 131 or MCDB 107 contg. isolated serum albumin, transferrin, hydrocortisone and epithelial growth factor.

ST vascular endothelial cell culture medium; polymer peptide adhesive cell

IT Blood vessel

(endothelium; method for serum-free culture of human vascular endothelial cells) IT Cell Culture media (method for serum-free culture of human vascular endothelial cells) Albumins, biological studies TT Polymers, biological studies Transferrins RL: BSU (Biological study, unclassified); BIOL (Biological study) (method for serum-free culture of human vascular endothelial cells) 50-23-7, Hydrocortisone 50-89-5, Thymidine, biological studies ΙT 50-99-7, D-Glucose, biological studies 52-90-4, Cysteine , biological studies 56-40-6, Glycine, biological studies 56-41-7, 56-45-1, L-Serine, biological studies Alanine, biological studies 56-85-9, Glutamine, 56-84-8, L-Aspartic acid, biological studies 56-86-0, Glutamic acid, biological studies 56-87-1, biological studies 58-05-9, Folinic acid 58-85-5, D-Biotin Lysine, biological studies 59-43-8, Thiamine, biological studies 60-18-4, Tyrosine, biological 63-68-3, Methionine, 61-90-5, L-Leucine, biological studies studies biological studies 63-91-2, Phenylalanine, biological studies 65-23-6, 70-47-3, Asparagine, 67-48-1 68-19-9, Vitamin B12 Pyridoxine 71-00-1, Histidine, biological studies 72 - 18 - 4. biological studies Valine, biological studies 72-19-5, Threonine, biological studies 73-22-3, Tryptophan, biological studies 73-24-5, Adenine, biological 73-32-5, Isoleucine, biological studies 74-79-3, L-Arginine, biological studies 79-83-4, D-Pantothenic acid 83-88-5, Riboflavin, biological studies 87-67-2, Choline bitartrate, biological studies 87-89-8, myo-Inositol 98-92-0, Nicotinamide 110-60-1, Putrescine 113-24-6, Sodium pyruvate 143-74-8, Phenol red 144-55-8, Carbonic acid monosodium salt, 137-08-6 147-85-3, Proline, biological studies 1200-22-2, biological studies 1344-09-8 1492-18-8, Calcium folinate .alpha.-Lipoic acid 7447-40-7, Potassium chloride (KCl), biological 7487-88-9, Sulfuric acid magnesium salt (1:1), studies biological studies 7558-79-4 7647-14-5, Sodium chloride, biological studies 7718-54-9, Nickel chloride (NiCl2), biological studies 7720-78-7, Sulfuric acid **iron**(2+) 7758-98-7, Sulfuric acid copper(2+) salt (1:1), 7733-02-0 salt (1:1) 7772-99-8, Tin chloride (SnCl2), biological biological studies 7778-77-0 7785-87-7 7803-55-6 10043-52-4, Calcium studies chloride (CaCl2), biological studies 10102-18-8 12027-67-7 110590-64-2 129058-83-9 134580-64-6 91037-65-9 93674-99-8 RL: BSU (Biological studý, unclassified); BIOL (Biological study) (method for serum-free culture of human vascular endothelial cells) ΙT 50-89-5, Thymidine, biological studies 52-90-4, Cysteine, biological studies 79-83-4, D-Pantothenic acid RL: BSU (Biological study, unclassified); BIOL (Biological study) (method for serum-free culture of human vascular endothelial cells) RN 50-89-5 HCAPLUS (CA INDEX NAME) Thymidine (8CI, 9CI) CN

Absolute stereochemistry.



RN 52-90-4 HCAPLUS

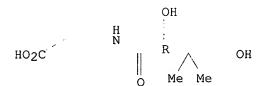
CN L-Cysteine (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 79-83-4 HCAPLUS

CN .beta.-Alanine, N-[(2R)-2,4-dihydroxy-3,3-dimethyl-1-oxobutyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).



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L100 ANSWER 9 OF 15 HCAPLUS COPYRIGHT 2003 ACS
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AN 1997:625591 HCAPLUS

DN 127:290229

TI Hematopoietic cell culture nutrient supplement

IN Daley, John P.; Dadey, Barbara M.; Biddle, William; Wysocki, Michelle G.

PA Life Technologies, Inc., USA

SO PCT Int. Appl., 73 pp.

CODEN: PIXXD2

DT Patent

LA English

IC ICM C12N005-00 ICS A61K035-28

CC 9-11 (Biochemical Methods)
 Section cross-reference(s): 13

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 9733978 A1 19970918 WO 1997-US1867 19970131

W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

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RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR,
             IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML,
             MR, NE, SN, TD, TG
                            19970918
                                            CA 1997-2248142 19970131
     CA 2248142
                       AΑ
     AU 9722600
                       A1
                            19971001
                                           AU 1997-22600
                                                             19970131
                            19990120
                                                             19970131
     EP 891419
                                           EP 1997-905789
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            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, FI
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                                            JP 1997-532595
                                                             19970131
                                           US 1997-792299
     US 2001033835
                       A1
                            20011025
                                                             19970131
PRAI US 1996-13149P
                       Р
                            19960312
     WO 1997-US1867
                       W
                            19970131
AB
     The present invention provides a serum-free supplement
     which supports the growth of hematopoietic cells in culture.
     supplement contains .gtoreq.1 ingredients selected from the group
     consisting of .gtoreq.1 antioxidant, .gtoreq.1 albumin or albumin
     substitute, .gtoreq.1 lipid agent, .gtoreq.1 insulin or insulin
     substitute, .gtoreq.1 transferrin or transferrin substitute, .gtoreq.1
     trace element, and .gtoreq.1 glucocorticoid, wherein a basal cell culture
    medium supplemented with the supplement is capable of supporting the
     expansion of CD34+ hematopoietic cells and cells of myeloid lineage, in
     serum-free culture. The present invention also provides
     methods for culturing and for differentiating hematopoietic cells.
ST
     hematopoietic cell culture serum free supplement;
    myeloid cell culture serum free supplement
IT
     Bone marrow
        (cells; hematopoietic cell culture nutrient supplement)
TΤ
     Hematopoietic precursor cell
        (erythroid burst-forming; hematopoietic cell culture nutrient
        supplement)
ΙT
     Cytometry
        (flow; hematopoietic cell culture nutrient supplement)
IT
     Animal tissue culture
     Antioxidants
     Blood serum
     Cell proliferation
     Hematopoiesis
     Hematopoietic precursor cell
        (hematopoietic cell culture nutrient supplement)
IT
     Acid-base indicators
     Albumins, biological studies
    Amino acids, biological studies
    Ape
     Buffers
     Cat (Felis catus)
     Cattle
     Cell differentiation
     Dog (Canis familiaris)
     Glucocorticoids
     Goat
     Growth factors, animal
     Guinea pig (Cavia porcellus)
     Hamster
     Horse (Equus caballus)
     Interleukin 3
     Lipids, biological studies
     Mammal (Mammalia)
     Monkey
     Mouse
     Nucleic acids
     Rabbit
     Rat
     Salts, biological studies
```

Sheep Stem cell factor Swine Trace elements, biological studies Transferrins Vitamins RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (hematopoietic cell culture nutrient supplement) IT Hematopoietic precursor cell (myeloid; hematopoietic cell culture nutrient supplement) ΙT Albumins, biological studies RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (serum; hematopoietic cell culture nutrient supplement) 50-23-7, Hydrocortisone 50-99-7, D-Glucose, biological studies TΤ 51-35-4, L-Hydroxyproline 52-90-4, L-Cysteine, biological studies 56-40-6, Glycine, biological studies 56-41-7, L-Alanine, biological studies 56-45-1, L-Serine, biological studies 56-84-8, L-Aspartic acid, biological studies 56-85-9, L-Glutamine, 56-86-0, L-Glutamic acid, biological studies biological studies 56-87-1, L-Lysine, biological studies 58-85-5, Biotin 59-30-3, Folic acid, biological studies 59-43-8, Thiamin, biological studies L-Tyrosine, biological studies 60-24-2, 2-Mercaptoethanol 61-90-5, L-Leucine, biological studies 63-68-3, L-Methionine, biological studies 63-91-2, L-Phenylalanine, biological studies 66-72-8, Pyridoxal 68-19-9, Vitamin B12 67-48-1, Choline chloride 70-47-3, L-Asparagine, biological studies 71-00-1, L-Histidine, 72-18-4, L-Valine, biological studies 72-19-5, biological studies L-Threonine, biological studies 73-22-3, L-Tryptophan, biological 73-32-5, L-Isoleucine, biological studies 74-79-3, L-Arginine, 83-88-5, Riboflavin, biological studies 87-89-8, biological studies i-Inositol 98-92-0, Niacinamide 113-24-6, **Sodium** pyruvate 137-08-6, Calcium D-127-17-3D, Pyruvic acid, salts 141-43-5, biological studies 143-74-8, Phenol red pantothenate 144-55-8, Carbonic acid monosodium salt, biological studies L-Proline, biological studies 616-91-1, N-Acetyl-L-cysteine 3812-32-6, Carbonate, biological 3812-32-6D, Carbonate, salts, biological studies 7365-45-9, HEPES 7439-95-4, Magnesium, biological studies 7439-95-4D, Magnesium, salts, biological studies 7440-09-7, Potassium, biological studies 7440-09-7D, Potassium, salts, biological studies 7440-23-5, Sodium, biological studies 7440-23-5D , Sodium, salts, biological studies 7440-70-2, Calcium, biological studies 7440-70-2D, Calcium 7447-40-7, Potassium , salts, biological studies chloride, biological studies 7487-88-9, Magnesium sulfate, biological studies 7558-80-7, Sodium dihydrogen 7647-14-5, Sodium chloride (NaCl), phosphate 7757-79-1, Potassium nitrate, biological biological studies 8049-62-5, Zinc 7782-49-2, Selenium, biological studies studies 9004-10-8, Insulin, biological studies 10043-52-4, insulin Calcium chloride (CaCl2), biological studies 11096-26-7, Erythropoietin 10102-18-8, Sodium selenite 14265-44-2, Phosphate, biological studies 14265-44-2D, Phosphate, salts, biological studies 83869-56-1, Granulocyte/macrophage colony-stimulating factor 52225-20-4 143011-72-7, Granulocyte-colony-stimulating factor RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BUU (Biological use, unclassified); BIOL (Biological

```
study); USES (Uses)
        (hematopoietic cell culture nutrient supplement)
     197179-47-8, Human Ex-Cyte
IT
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
     study); USES (Uses)
        (human; hematopoietic cell culture nutrient supplement)
ΙT
     52-90-4, L-Cysteine, biological studies 616-91-1
     , N-Acetyl-L-cysteine 7439-95-4,
    Magnesium, biological studies 7439-95-4D,
    Magnesium, salts, biological studies 7440-09-7,
     Potassium, biological studies 7440-09-7D,
     Potassium, salts, biological studies 7440-23-5,
     Sodium, biological studies 7440-23-5D, Sodium,
     salts, biological studies 7440-70-2, Calcium,
     biological studies 7440-70-2D, Calcium, salts,
     biological studies 14265-44-2, Phosphate, biological
     studies 14265-44-2D, Phosphate, salts, biological
     studies
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
     study); USES (Uses)
        (hematopoietic cell culture nutrient supplement)
     52-90-4 HCAPLUS
RN
CN
     L-Cysteine (9CI) (CA INDEX NAME)
Absolute stereochemistry.
      NH<sub>2</sub>
            SH
HO<sub>2</sub>C
RN
     616-91-1 HCAPLUS
CN
     L-Cysteine, N-acetyl- (9CI) (CA INDEX NAME)
Absolute stereochemistry.
     R CO2H
       NHAc
RN
     7439-95-4 HCAPLUS
CN
     Magnesium (8CI, 9CI)
                           (CA INDEX NAME)
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Mg 7439-95-4 HCAPLUS RN CN Magnesium (8CI, 9CI) (CA INDEX NAME) Mg RN 7440-09-7 HCAPLUS CN Potassium (8CI, 9CI) (CA INDEX NAME) K

RN 7440-09-7 HCAPLUS

CN Potassium (8CI, 9CI) (CA INDEX NAME)

K

RN 7440-23-5 HCAPLUS

CN Sodium (8CI, 9CI) (CA INDEX NAME)

Na

RN 7440-23-5 HCAPLUS

CN Sodium (8CI, 9CI) (CA INDEX NAME)

Na

RN 7440-70-2 HCAPLUS

CN Calcium (8CI, 9CI) (CA INDEX NAME)

Ca

RN 7440-70-2 HCAPLUS

CN Calcium (8CI, 9CI) (CA INDEX NAME)

Ca

RN 14265-44-2 HCAPLUS

CN Phosphate (8CI, 9CI) (CA INDEX NAME)

RN 14265-44-2 HCAPLUS

CN Phosphate (8CI, 9CI) (CA INDEX NAME)

L100 ANSWER 10 OF 15 HCAPLUS COPYRIGHT 2003 ACS AN 1997:483452 HCAPLUS

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127:92426
DN
ΤI
     Serum-free culture medium for Drosophila insect cells
     Ramos, Luciano; Murnane, Amy Anne; Oka, Melvin Susumu
ΤN
PΑ
     SmithKline Beecham Corp., USA
     U.S., 5 pp., Cont. of U.S. Ser. No. 183,585, abandoned.
SO
     CODEN: USXXAM
DT
     Patent
LA
    English
IC
     ICM C12N005-00
NCL
    435404000
CC
     9-11 (Biochemical Methods)
     Section cross-reference(s): 12
FAN.CNT 1
                     KIND DATE
                                           APPLICATION NO.
     PATENT NO.
                                                            DATE
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                           -----
                                           -----
    US 5641678
                                                            19950607
                            19970624
                                           US 1995-483634
PΙ
                      Α
                            19940118
PRAI US 1994-183585
     The serum-free medium of the invention comprises a
    basal medium to which is added yeast hydrolyzate and albumin or dextran.
     In another embodiment of the invention, albumin hydrolyzate is added to
     the basal medium, in addn. to the aforementioned compds.
     Drosophila cell culture serum free medium; insect cell
ST
     culture serum free media
    Protein hydrolyzates
IΤ
    Protein hydrolyzates
    RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (albumin hydrolyzates; serum-free culture medium
        for Drosophila insect cells)
    Envelope proteins
TT
     RL: BSU (Biological study, unclassified); MFM (Metabolic formation); BIOL
     (Biological study); FORM (Formation, nonpreparative)
        (gp120env; serum-free culture medium for Drosophila
       insect cells)
TΤ
    Yeast
        (hydrolyzate; serum-free culture medium for
       Drosophila insect cells)
TΤ
    Albumins, biological studies
    Albumins, biological studies
    Lactalbumins
    Lactalbumins
    RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (hydrolyzates; serum-free culture medium for
       Drosophila insect cells)
    Protein hydrolyzates
TT
     Protein hydrolyzates
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (lactalbumin hydrolyzates; serum-free culture
       medium for Drosophila insect cells)
    Animal tissue culture
TT
    Culture media
    Drosophila
     Drosophila melanogaster
     Insect (Insecta)
        (serum-free culture medium for Drosophila insect
        cells)
IT
     Albumins, biological studies
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (serum-free culture medium for Drosophila insect
       cells)
```

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IT
     Albumins, biological studies
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (serum; serum-free culture medium for
        Drosophila insect cells)
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ΙT
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     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
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        cells)
     52-90-4, L-Cysteine, biological studies
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     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (serum-free culture medium for Drosophila insect
        cells)
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RN
     52-90-4
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                      (CA INDEX NAME)
CN
Absolute stereochemistry.
      NH2
            SH
      R
HO<sub>2</sub>C
L100 ANSWER 11 OF 15 HCAPLUS COPYRIGHT 2003 ACS
AN
     1997:454000 HCAPLUS
DN
     127:62876
     Immortalized human skin cell lines and serum-free
TΤ
     medium for their culture
     Baur, Markus; Mace, Catherine; Malnoe, Armand; Pfeifer, Andrea M. A.;
IN
     Regnier, Marcelle
PΑ
     Societe Des Produits Nestle S.A., Switz.
SO
     Eur. Pat. Appl., 27 pp.
     CODEN: EPXXDW
DT
     Patent
LA
     French
IC
     ICM C12N005-08
     ICS C12N005-00; C12N005-22
CC
     9-11 (Biochemical Methods)
     Section cross-reference(s): 13, 14
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                                           APPLICATION NO.
                                                            DATE
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PΙ
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                                                            19961219
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                      A1
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             SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM
         RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR,
             IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML,
            MR, NE, SN, TD, TG
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                                           ES 1996-203641
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PRAI US 1995-576483
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                            19961219
    WO 1996-EP5818
                      W
                            19961219
     US 1998-91483
                      A1
                            19980619
AΒ
     The invention concerns immortalized cell lines, esp. of keratinocytes and
    melanocytes derived from normal human skin, as well as a novel
     serum-free medium for the isolation, growth,
     and maintenance of these cells. Procedures and compns. are disclosed for
     producing primary melanocytes and keratinocytes in the absence of serum
     and without fibroblast nurse cells. Plasmids derived from SV40 virus or
     papilloma virus 16 were used to immortalize the melanocytes and
     keratinocytes of this invention. The findings are useful for the improved
     immunol., pharmacol., photo-, and chemotoxicol. anal. of cutaneous
     reactions and for the expression of heterologous genes.
                                                              The cells may be
     used for studying the inflammation reaction and for skin grafting.
ST
     immortalized keratinocyte melanocyte cell culture medium
     ; skin graft immortalized keratinocyte melanocyte; inflammation analysis
     immortalized keratinocyte melanocyte
IT
    Animal cell line
        (DK2-NR; immortalized human skin cell lines culture in
        serum-free medium)
ΙT
    Animal cell line
        (DK3-NR; immortalized human skin cell lines culture in
        serum-free medium)
IT
     Animal cell line
        (DM2-NR; immortalized human skin cell lines culture in
        serum-free medium)
IT
     Animal cell line
        (FK2-NR; immortalized human skin cell lines culture in
       serum-free medium)
IT
     Reagents
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (MCDB 153; immortalized human skin cell lines culture in
```

```
serum-free medium)
TΤ
    Reagents
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (NR-2; immortalized human skin cell lines culture in
        serum-free medium)
IT
    Reagents
    RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (NR-3; immortalized human skin cell lines culture in
        serum-free medium)
IT
     Reagents
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (NR-4; immortalized human skin cell lines culture in
        serum-free medium)
    Fatty acids, biological studies
TΤ
    RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (essential; immortalized human skin cell lines culture in
       serum-free medium)
ΙT
    Gene
        (expression; immortalized human skin cell lines culture in
       serum-free medium)
ΙT
    Transformation, neoplastic
        (immortalization; immortalized human skin cell lines culture
       in serum-free medium)
    Animal tissue culture
TT
    Human papillomavirus 16
    Inflammation
    Melanocyte
    Physiological saline solutions
    Plasmids
    Simian virus 40
    Skin
        (immortalized human skin cell lines culture in serum
       -free medium)
ΙT
    Amino acids, biological studies
    Antibiotics
    Pituitary hormones
    Salts, biological studies
    Transferrins
    Vitamins
    RL: BAC (Biological activity or effector, except adverse); BSU (Biological
    study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
    study); USES (Uses)
        (immortalized human skin cell lines culture in serum
        -free medium)
IT
    Filaggrin
    RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); BIOL (Biological study); FORM (Formation,
    nonpreparative); OCCU (Occurrence)
        (immortalized human skin cell lines culture in serum
        -free medium)
ΙT
    Keratins
    RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); BIOL (Biological study); FORM (Formation,
    nonpreparative); OCCU (Occurrence)
        (immortalized human skin cell lines culture in serum
        -free medium)
ΙT
    Melanins
    RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
```

(Metabolic formation); BIOL (Biological study); FORM (Formation,

```
nonpreparative); OCCU (Occurrence)
        (immortalized human skin cell lines culture in serum
        -free medium)
IT
     Proteins, general, biological studies
     RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); BIOL (Biological study); FORM (Formation,
     nonpreparative); OCCU (Occurrence)
        (immortalized human skin cell lines culture in serum
        -free medium)
IT
     Tumor necrosis factors
     RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); BIOL (Biological study); FORM (Formation,
     nonpreparative); OCCU (Occurrence)
        (immortalized human skin cell lines culture in serum
        -free medium)
IΤ
     Fibronectins
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (immortalized human skin cell lines culture in serum
        -free medium)
TΤ
     Proteins, specific or class
     RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); BIOL (Biological study); FORM (Formation,
     nonpreparative); OCCU (Occurrence)
        (involucrins; immortalized human skin cell lines culture in
        serum-free medium)
TΤ
    Skin
        (keratinocyte; immortalized human skin cell lines culture in
        serum-free medium)
     Proteins, specific or class
IT
    RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); BIOL (Biological study); FORM (Formation,
     nonpreparative); OCCU (Occurrence)
        (loricrins; immortalized human skin cell lines culture in
        serum-free medium)
    Amino acids, biological studies
IT
    RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
     study); USES (Uses)
        (salts; immortalized human skin cell lines culture in
        serum-free medium)
TT
    Transplant and Transplantation
        (skin; immortalized human skin cell lines culture in
        serum-free medium)
IT
    Skin
        (transplant; immortalized human skin cell lines culture in
        serum-free medium)
IT
     50-23-7, Hydrocortisone 50-89-5, Thymidine, biological studies
     50-99-7, D-Glucose, biological studies
                                              51-43-4, Epinephrine
    52-90-4, L-Cysteine, biological studies
                                              56-40-6,
    Glycine, biological studies
                                  56-41-7, L-Alanine, biological studies
     56-45-1, L-Serine, biological studies
                                             56-84-8, L-Aspartic acid,
                          56-85-9, L-Glutamine, biological studies
    biological studies
                                                                      56-86-0,
    L-Glutamic acid, biological studies
                                          56-87-1, L-Lysine, biological
               57-92-1, Streptomycin, biological studies
                                                           58-85-5
                                                                     59-30-3,
                                    59-43-8, Thiamin, biological studies
     Folic acid, biological studies
     60-18-4, L-Tyrosine, biological studies
                                             61-33-6, biological studies
     61-90-5, L-Leucine, biological studies
                                              63-68-3, L-Methionine, biological
     studies
               63-91-2, L-Phenylalanine, biological studies
                                                              65-23-6,
                                              68-19-9,
     Pyridoxine
                  67-48-1, Choline chloride
    Cyanocobalamin
                      70-47-3, L-Asparagine, biological studies
                                                                  71-00-1,
    L-Histidine, biological studies
                                      72-18-4, L-Valine, biological studies
```

73-22-3, L-Tryptophan,

72-19-5, L-Threonine, biological studies

```
biological studies
                      73-24-5, Adenine, biological studies
                                                             73-32-5.
 L-Isoleucine, biological studies 74-79-3, L-Arginine, biological studies
 83-88-5, Riboflavin, biological studies
                                           87-89-8, i-Inositol
 Nicotinamide
                110-60-1, Putrescine
                                      113-24-6, Sodium pyruvate
                            137-08-6, Calcium
 127-09-3, Sodium acetate
                141-43-5, biological studies
                                               143-74-8, Phenol red
 pantothenate
 144-55-8, Carbonic acid monosodium salt, biological studies
                                                               147-85-3,
 L-Proline, biological studies
                               1071-23-4, Phosphoethanolamine
 1077-28-7, Thioctic acid
                           1397-89-3, Fungizone
                                                   6834-92-0
                                                               7365-45-9,
 HEPES 7440-70-2, Calcium, biological studies
 7447-40-7, Potassium chloride, biological studies
 7558-79-4, Disodium phosphate
                                 7647-14-5, Sodium
 chloride (NaCl), biological studies
                                       7733-02-0, Zinc sulfate
 7758-98-7, Cupric sulfate, biological studies
                                                 7772-99-8, Tin
 chloride, biological studies
                                7773-01-5, Manganese
 chloride
           7786-30-3, Magnesium chloride,
                      7786-81-4, Nickel sulfate
                                                  7803-55-6, Ammonium
 biological studies
                9004-10-8, Insulin, biological studies 10028-22-5, Ferric
 metavanadate
           10043-52-4, -Calcium chloride (CaCl2),
 biological studies
                      10102-18-8, Sodium selenite
                                                    12027-67-7,
 Ammonium molybdate
                      16561-29-8, Phorbol 12-myristate 13-acetate
                                      106096-93-9, Basic fibroblast growth
 62229-50-9, Epidermal growth factor
 factor
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological
 study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
 study); USES (Uses)
    (immortalized human skin cell lines culture in serum
    -free medium)
 50812-37-8, Glutathione S-transferase
 RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
 (Metabolic formation); BIOL (Biological study); FORM (Formation,
 nonpreparative); OCCU (Occurrence)
    (immortalized human skin cell lines culture in serum
    -free medium)
 9035-51-2, Cytochrome P 450, biological studies
 RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
 (Metabolic formation); BIOL (Biological study); FORM (Formation,
 nonpreparative); OCCU (Occurrence)
    (multiple forms; immortalized human skin cell lines culture
    in serum-free medium)
 9001-12-1, Collagenase
 RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
 (Metabolic formation); BIOL (Biological study); FORM (Formation,

    nonpreparative); OCCU (Occurrence)

    (type I; immortalized human skin cell lines culture in
    serum-free medium)
 50-89-5, Thymidine, biological studies 52-90-4, L-
 Cysteine, biological studies 7440-70-2, Calcium
 , biological studies
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological
 study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
 study); USES (Uses)
    (immortalized human skin cell lines culture in serum
    -free medium)
 50-89-5 HCAPLUS
                      (CA INDEX NAME)
 Thymidine (8CI, 9CI)
```

Absolute stereochemistry.

TT

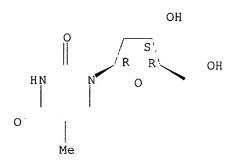
TΤ

IT

TT

RN

CN



RN 52-90-4 HCAPLUS CN L-Cysteine (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 7440-70-2 HCAPLUS CN Calcium (8CI, 9CI) (CA INDEX NAME)

Ca

L100 ANSWER 12 OF 15 HCAPLUS COPYRIGHT 2003 ACS

AN 1997:16735 HCAPLUS

DN 126:141671

TI Development of a chemically defined **medium** for the growth of Leuconostoc mesenteroides

AU Foucaud, Catherine; Francois, Alan; Richard, Jean

CS Unite de Recherches Laitieres, Institut National de la Recherche Agronomique, Jouy en Josas, 78350, Fr.

SO Applied and Environmental Microbiology (1997), 63(1), 301-304 CODEN: AEMIDF; ISSN: 0099-2240

PB American Society for Microbiology

DT Journal

LA English

CC 9-11 (Biochemical Methods)

AB A chem. defined medium for the growth of Leuconostoc mesenteroides was developed. This medium contained lactose, Mn2+, Mg2+, 12 amino acids, eight vitamins, adenine, uracil, and Tween 80. We showed the beneficial effect of aerobic conditions on growth and that potassium phosphate (135 mM) is a suitable buffer. The growth rate in this medium was 0.85 .+-. 0.10 h-1 for the six strains examd., and cell densities up to 3.5 .times. 109 CFU/mL were attained.

ST Leuconostoc growth chem defined medium

IT Culture media

Leuconostoc mesenteroides

(development of a chem. defined **medium** for the growth of Leuconostoc mesenteroides)

IT Amino acids, biological studies
 Vitamins

RL: BUU (Biological use, unclassified); BIOL (Biological study); USES

```
(Uses)
        (development of a chem. defined medium for the growth of
       Leuconostoc mesenteroides)
ΙT
    50-89-5, Thymidine, biological studies 52-90-4, L-
    Cysteine, biological studies 56-40-6, Glycine, biological studies 56-41-7, L-Alanine, biological studies 56-45-1, L-Serine,
    biological studies
                         56-84-8, L-Aspartic acid, biological studies
    56-85-9, L-Glutamine, biological studies 56-87-1, L-Lysine, biological
               58-63-9, Inosine 58-85-5, Biotin
                                                    59-30-3, Folic acid,
                        59-67-6, Nicotinic acid, biological studies
    biological studies
     60-18-4, L-Tyrosine, biological studies
                                               61-90-5, Leu, biological studies
    63-42-3, Lactose 63-68-3, L-Methionine, biological studies
                                                                   63-91-2,
    L-Phenylalanine, biological studies
                                          65-22-5
                                                    65-86-1, Orotic acid
     66-22-8, Uracil, biological studies
                                           67-03-8, Thiamine dichloride
    68-19-9, Vitamin B12
                            69-89-6, Xanthine
                                                70-47-3, Asn, biological
             71-00-1, L-Histidine, biological studies
                                                          72-18-4, L-Valine,
    studies
                        72-19-5, L-Threonine, biological studies
    biological studies
                                                                     73-22-3,
    L-Tryptophan, biological studies 73-24-5, Adenine, biological studies
    73-32-5, L-Isoleucine, biological studies 73-40-5, Guanine
                                                                   74-79-3,
    L-Arginine, biological studies 83-88-5, Riboflavin, biological studies
    127-09-3, Sodium acetate 137-08-6, Calcium
    pantothenate 147-85-3, L-Proline, biological studies
                                                              524-36-7
    555-06-6, Sodium p-aminobenzoate
                                        1077-28-7, DL-6,8-Thioctic
    acid
            3458-72-8
                        7646-79-9, Cobalt chloride (CoCl2),
    biological studies 7733-02-0 7758-11-4
                                                 7758-98-7, Sulfuric acid
    copper(2+) salt (1:1), biological studies
                                                 7778-77-0 7785-87-7
    7786-30-3, Magnesium chloride (MgCl2), biological
              9005-65-6, Tween 80. 10043-52-4, Calcium
    studies
    chloride (CaCl2), biological studies
                                           12040-57-2, Iron
               16397-91-4, Mn2+, biological studies 22537-22-0
    chloride
     , Mg2+, biological studies
    RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (development of a chem. defined medium for the growth of
       Leuconostoc mesenteroides)
ΙT
    50-89-5, Thymidine, biological studies 52-90-4, L-
    Cysteine, biological studies 22537-22-0, Mg2+,
    biological studies
    RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (development of a chem. defined medium for the growth of
```

(CA INDEX NAME)

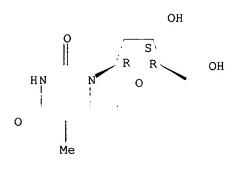
Absolute stereochemistry.

50-89-5 HCAPLUS

Thymidine (8CI, 9CI)

RN

CN



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RN 52-90-4 HCAPLUS
CN L-Cysteine (9CI) (CA INDEX NAME)
```

Leuconostoc mesenteroides)

for)

```
Absolute stereochemistry.
      VH2
            SH
     R
HO<sub>2</sub>C
RN
    22537-22-0 HCAPLUS
    Magnesium, ion (Mg2+) (8CI, 9CI) (CA INDEX NAME)
CN
Mq^{2}+
L100 ANSWER 13 OF 15 HCAPLUS COPYRIGHT 2003 ACS
AN
    1994:72906 HCAPLUS
DN
    120:72906
ΤI
    Serum-free culture medium for eukaryotic cells
IN
    Bosslet, Klaus
PΑ
    Behringwerke AG, Germany
SO
    Eur. Pat. Appl., 7 pp.
    CODEN: EPXXDW
DT
    Patent
LA
    German
    ICM C12N005-00
TC
    9-11 (Biochemical Methods)
CC
FAN.CNT 1
    PATENT NO.
                     KIND DATE
                                         APPLICATION NO. DATE
    _____
                     ----
                                          _____
                                     EP 1993-107948 19930515
    EP 573812
                    A2 19931215
PΙ
       R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE
    DE 4219250 A1 19931216
                                          DE 1992-4219250 19920612
    CA 2098255
                      AA
                           19931213
                                          CA 1993-2098255
                                                          19930611
    AU 9340183
                      A1
                           19931216
                                          AU 1993-40183
                                                           19930611
                     A2
                                          JP 1993-140227
    JP 06078759
                           19940322
                                                           19930611
PRAI DE 1992-4219250
                           19920612
    A culture medium is described which contains amino acids, vitamins,
    cofactors, inorg. salts, ethanolamine, Na2SeO3, human transferrin, and
    human insulin, and is otherwise free of animal proteins. This medium
    supports the growth of a variety of eukaryotic cells (e.g. human and rat
    tumor cells) as well as media contg. fetal calf serum, and is much less
    expensive.
ST
    ethanolamine culture medium animal cell; selenite culture medium animal
    cell; transferrin culture medium animal cell; insulin culture medium
    animal cell
IT
    Amino acids, biological studies
    Coenzymes
    Salts, biological studies
    Vitamins
    RL: BIOL (Biological study)
        (culture medium for animal cells contg.)
    Transferrins
IT
    RL: BIOL (Biological study)
        (culture medium for animal cells contg. human)
IT
    Animal tissue culture
        (ethanolamine and insulin and selenite and transferrin in growth medium
```

IT 50-99-7, Glucose, biological studies 52-90-4, Cysteine , biological studies 56-40-6, Glycine, biological studies Alanine, biological studies 56-45-1, Serine, biological studies 56-84-8, Aspartic acid, biological studies 56-85-9, Glutamine,

biological studies 56-86-0, Glutamic acid, biological studies 59-30-3, Folic acid, biological studies 60-18-4, Tyrosine, biological studies 61-90-5, L-Leucine, biological studies 63-68-3, Methionine, biological 63-91-2, Phenylalanine, biological studies 65-22-5, Pyridoxal hydrochloride 67-03-8, Thiamine hydrochloride 67-48-1, Choline 70-47-3, Asparagine, biological studies 72 - 18 - 4, chloride Valine, biological studies 72-19-5, Threonine, biological studies 73-22-3, Tryptophan, biological studies 73-32-5, Isoleucine, biological 83-88-5, Riboflavin, biological studies 87-89-8, Inositol 113-24-6, **Sodium** pyruvate 137-08-6, 98-92-0, Nicotinamide Calcium pantothenate 141-43-5, biological studies 143-74-8, Phenol red 144-55-8, Sodium bicarbonate, biological 147-85-3, Proline, biological studies 645-35-2, Histidine 657-27-2, Lysine hydrochloride 1119-34-2, Arginine hydrochloride 7447-40-7, Potassium chloride (KCl), hydrochloride biological studies 7487-88-9, Magnesium sulfate, biological 7558-80-7, Sodium dihydrogen phosphate 7647-14-5, Sodium chloride, biological studies 10043-52-4, Calcium chloride, biological studies 10102-18-8, Sodium selenite 10421-48-4, Ferric nitrate RL: BIOL (Biological study) (culture medium for animal cells contg.) 9004-10-8, Insulin, biological studies ΙT RL: BIOL (Biological study) (culture medium for animal cells contg. human) TΤ 52-90-4, Cysteine, biological studies RL: BIOL (Biological study) (culture medium for animal cells contg.) 52-90-4 HCAPLUS RN CN L-Cysteine (9CI) (CA INDEX NAME)

Absolute stereochemistry.

```
L100 ANSWER 14 OF 15 HCAPLUS COPYRIGHT 2003 ACS
     1990:32876 HCAPLUS
ΑN
DN
     112:32876
ΤI
     Improved maintenance of adult rat hepatocytes in a new serum-
     free medium in the presence or absence of barbiturates
ΑU
    Miyazaki, Masahiro; Suzuki, Yasunori; Oda, Munehiro; Kawai, Akira; Bai,
    Liyan; Sato, Jiro
CS
    Med. Sch., Okayama Univ., Okayama, 700, Japan
     In Vitro Cellular & Developmental Biology (1989), 25(9), 839-48
SO
     CODEN: ICDBEO; ISSN: 0883-8364
DT
     Journal
LA
     English
CC
     9-11 (Biochemical Methods)
     Section cross-reference(s): 1, 13
     For serum-free primary culture of adult rat
AB
```

hepatocytes, a synthetic medium DM-160 and rat-tail collagen were selected for the basal medium and for the culture substratum, resp. Barbiturates, such as phenobarbital and 1-ethyl-5-isobutylbarbiturate, efficiently supported survival of hepatocytes and maintained their morphol. features at lower concns. under the serum-free conditions than under the serum-supplemented conditions. However, the hepatocyte survival rates under the serum-free conditions were lower than those under the serum-supplemented conditions in the presence or absence

of barbiturates. Supplementation of the basal medium with a combination of 5 groups of factors (5Fs), such as 8 amino acids (Ala, Arg, Gly, Ile, Met, Phe, Pro, and Trp), 2 unsatd. fatty acids (linoleate and oleate), protease inhibitor (aprotinin), 3 vitamins (A, C, and E), and 5 trace elements (Mn, Fe, Cu, Zn, and Se), improved hepatocyte survival under the serum-free conditions in the presence or absence of barbiturates. In other words, the serum could be completely substituted by the 5Fs. Hepatocyte cultures maintained in the 5Fs-supplemented basal medium showed excellent induction of tyrosine aminotransferase activity in response to dexamethasone in the presence or absence of barbiturates. The efficiency of the 5Fs-supplemented basal medium for maintaining hepatocytes was not inferior to those of other media in common use with hepatocytes, such as Williams' medium E and Waymouth's medium MB-752/1. In conclusion, maintenance of functional hepatocytes in serum-free primary culture could be improved by use of the new medium prepn. (the 5Fs-supplemented DM-160) in the presence of barbiturates. hepatocyte culture serum free media barbiturate Animal tissue culture

ST

TΤ

(of hepatocytes, serum-free medium for, in barbiturates absence and presence)

ΙT Amino acids, biological studies Trace elements, biological studies Vitamins

RL: BIOL (Biological study)

(serum-free medium contg., for hepatocyte culture, barbiturates in relation to)

TT Liver

> (hepatocyte, culture of, serum-free medium for, in barbiturates absence and presence)

109791-24-4 67-52-7D, 2,4,6(1H,3H,5H)-Pyrimidinetrione, derivs. IT 50-06-6, Phenobarbital, biological studies

RL: ANST (Analytical study)

(hepatocyte culture on serum-free medium in presence of)

ΙT 9014-55-5, Tyrosine aminotransferase

RL: ANST (Analytical study)

(of hepatocyte in culture with serum-free medium, dexamethasone induction of)

68-26-8, Vitamin A TT 68-19-9, Cyanocobalamine 70-47-3, L-Asparagine, 71-00-1, L-Histidine, biological studies biological studies L-Valine, biological studies 72-19-5, L-Threonine, biological studies 73-22-3, Tryptophan, biological studies 73-32-5, Isoleucine, biological studies 74-79-3, L-Arginine, biological studies 79-83-4 83-88-5, Riboflavine, biological studies 87-89-8, Inositol 98-92-0, 143-74-8, Phenol Nicotinamide 112-80-1, Oleic acid, biological studies 144-55-8, Carbonic acid monosodium salt, biological studies 147-85-3, Proline, biological studies 1406-18-4, Vitamin E 7439-89-6, Iron, biological studies 7439-96-5, 7440-50-8, Copper, biological studies Manganese, biological studies 7440-66-6, Zinc, biological studies 7447-40-7, Potassium 7487-88-9, Sulfuric acid chloride (KCl), biological studies magnesium salt (1:1), biological studies 7558-80-7 7647-14-5, Sodium chloride (NaCl), biological studies 7782-49-2, 10043-52-4, Selenium, biological studies 9087-70-1, Aprotinin Calcium chloride (CaCl2), biological studies 50-81-7, Vitamin C, biological studies 50-99-7, Glucose, biological studies 52-90-4, L-Cysteine, biological studies 56-40-6, Glycine, biological studies 56-41-7, L-Alanine, biological studies 56-45-1, L-Serine, biological studies 56-84-8, L-Aspartic acid, 56-85-9, L-Glutamine, biological studies biological studies L-Glutamic acid, biological studies 56-87-1, L-Lysine, biological 58-85-5, Biotin 59-30-3, Folic acid, biological studies studies

59-43-8, Thiamine, biological studies 60-18-4, L-Tyrosine, biological 60-33-3, 9,12-Octadecadienoic acid (Z,Z)-, biological studies studies 61-90-5, L-Leucine, biological studies **62-49-7** 63-68-3, Methionine, biological studies 63-91-2, L-Phenylalanine, biological 65-23-6, Pyridoxine RL: ANST (Analytical study) (serum-free medium contg., for hepatocyte culture, barbiturates in relation to) 50-02-2, Dexamethasone RL: ANST (Analytical study) (tyrosine aminotransferase of hepatocytes in culture in serum -free medium induction by) 79-83-4 7439-89-6, Iron, biological studies 52-90-4, L-Cysteine, biological studies 62-49-7 RL: ANST (Analytical study) (serum-free medium contg., for hepatocyte culture, barbiturates in relation to)

RN 79-83-4 HCAPLUS

CN .beta.-Alanine, N-[(2R)-2,4-dihydroxy-3,3-dimethyl-1-oxobutyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).

RN 7439-89-6 HCAPLUS CN Iron (7CI, 8CI, 9CI) (CA INDEX NAME)

Fe

ΙT

IT

RN 52-90-4 HCAPLUS CN L-Cysteine (9CI) (CA INDEX NAME)

Absolute stereochemistry.

NH<sub>2</sub>

NH<sub>2</sub>

SH

RN 62-49-7 HCAPLUS

CN Ethanaminium, 2-hydroxy-N,N,N-trimethyl- (9CI) (CA INDEX NAME)

Me3+N-CH2-CH2-OH

L100 ANSWER 15 OF 15 HCAPLUS COPYRIGHT 2003 ACS
AN 1989:532627 HCAPLUS
DN 111:132627
TI Culture medium composition for animal cell culture
PA Grace, W. R., and Co., USA
SO Jpn. Kokai Tokkyo Koho, 19 pp.

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CODEN: JKXXAF
DT
     Patent
     Japanese
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TC
     ICM C12N005-00
ICA
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     JP 63267269
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                                                            19880318
         R: DE, FR, GB, IT
PRAI US 1987-29577
                            19870324
     Basal nutrient medium suitable for serum-free/-low
     tissue culture, e.g. hybridoma culture for manufg. monoclonal antibodies
     (MAb), is disclosed. MAb-producing CRL 1606 hybridoma cells were cultured
     in a serum-free medium reconstructed from the dried
     medium formulation (mosm 295) for up to 144 h. Cellular growth of and MAb
     prodn. by the cells in the medium thus prepd. were more efficient than the
     control group where the cells were cultured in DME medium supplemented
     with 10% fetal calf serum. For example, the cell no. and MAb prodn. at
     120 h was 2300.times.109/L and 262.9 mg/L, resp., compared to
     2000.times.109 and 149.6 for the control.
     animal tissue culture medium compn; serum free medium
ST
     monoclonal antibody manuf
IT
     Mammal
        (cells of, cultivation of, serum-free/-low culture
        medium for)
TΤ
     Carbohydrates and Sugars, biological studies
     Coenzymes
     Nucleic acids
     Vitamins
     RL: BIOL (Biological study)
        (culture medium contg., serum-free/-low, for animal
        tissue culture)
ΙT
     Amino acids, biological studies
     Lipids, biological studies
     Trace elements, biological studies
     RL: BIOL (Biological study)
        (culture medium contg., serum-free/low, for animal
        tissue culture)
IΤ
     Blood serum
     Proteins, biological studies
     RL: BIOL (Biological study)
        (culture medium low in, for animal tissue culture)
IT
     Albumins, biological studies
     RL: BIOL (Biological study)
        (culture medium supplemented with, serum-free/low)
ΙT
     Transferrins
     RL: BIOL (Biological study)
        (iron-satd., culture medium supplemented with, serum
        -free/-low)
ΙT
     Animal tissue culture
        (medium formulation for, serum-free/-low)
ΙT
     рН
        (of serum-free/-low culture medium, for animal
        tissue culture)
     Amino acids, biological studies
IT
     RL: BIOL (Biological study)
        (essential, culture medium contg., serum-free/low,
        for animal tissue culture)
ΙT
     Antibodies
     RL: BMF (Bioindustrial manufacture); BIOL (Biological study); PREP
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(Preparation) (monoclonal, manuf. of, serum-free/-low culture medium in relation to) 9004-10-8, Insulin, biological studies IT RL: BIOL (Biological study) (culture medium supplemented with, serum-free/low) 50-99-7, D-Glucose, biological studies 51-35-4, L-Hydroxy proline IT 56-40-6, Glycine, biological studies 56-41-7, L-Alanine, biological 56-45-1, L-Serine, biological studies 56-84-8, L-Aspartic blogical studies 56-85-9, L-Glutamine, biological studies acid, biological studies 58-85-5, Biotin 58-56-0, Pyridoxine hydrochloride 59-30-3, Folic acid, biological studies 60-33-3, 9,12-Octadecadienoic acid (Z,Z)-, biological61-90-5, L-Leucine, biological studies 63-68-3, L-Methionine, biological studies 63-91-2, L-Phenylalanine, biological studies 65-22-5, Pyridoxal hydrochloride 67-03-8, Thiamine hydrochloride 67-48-1, Choline chloride 68-19-9, Vitamin B12 70-18-8, Glutathione, biological studies 72-18-4, L-Valine, biological studies 72-19-5, L-Threonine, biological studies 73-22-3, 73-24-5, Adenine, biological studies L-Tryptophane, biological studies 73-32-5, L-Isoleucine, biological studies 74-79-3, L-Arginine, 83-88-5, Riboflavin, biological studies 87-89-8, biological studies i-Inositol 98-92-0, 3-Pyridinecarboxamide 113-24-6 137-08-6 147-85-3, 144-55-8, Sodium bicarbonate, biological studies 657-27-2, L-Lysine L-Proline, biological studies 333-93-7 1200-22-2, Lipoic acid 1310-73-2, Sodium hydrochloride hydroxide, biological studies 1321-11-5, Aminobenzoic acid 2002-24-6, Ethanolamine hydrochloride 5794-13-8, L-Asparagine monohydrate 5934-29-2, L-Histidine hydrochloride monohydrate 6035-45-6 7048-04-6, 7365-45-9, HEPES L-Cysteine hydrochloride, monohydrate 7446-20-0, Zinc sulfate (heptahydrate) 7447-40-7, Potassium 7647-14-5, Sodium chloride, biological studies 7758-99-8 7782-61-8 7782-63-0 chloride, biological studies 10101-97-0 12054-85-2 7782-85-6 7803-55-6 10034-99-8 10035-04-8 45738-97-4 26970-82-1 122666-87-9 13472-35-0 13517-24-3 122666-88-0, Thymidine hydrochloride RL: BIOL (Biological study) (medium formulation contg., for serum-low/-free mammalian cell culture) TΤ 12408-02-5 RL: BIOL (Biological study) (pH, of serum-free/-low culture medium, for animal tissue culture)

=> fil wpix FILE 'WPIX' ENTERED AT 17:33:14 ON 26 APR 2003 COPYRIGHT (C) 2003 THOMSON DERWENT

FILE LAST UPDATED: 16 APR 2003 <20030416/UP>
MOST RECENT DERWENT UPDATE: 200325 <200325/DW>
DERWENT WORLD PATENTS INDEX SUBSCRIBER FILE, COVERS 1963 TO DATE

Due to data production problems the WPI file had to be reset to update 200324. SDIs for update 24 will be rerun. The previous SDI run for 24 has been credited. Also answer sets created after April 10 may at least temporarily be affected and hence partially invalid.

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- >>> SLART (Simultaneous Left and Right Truncation) is now available in the /ABEX field. An additional search field

/BIX is also provided which comprises both /BI and /ABEX <<<

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    http://www.derwent.com/userguides/dwpi guide.html <<<
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L127 ANSWER 1 OF 2 WPIX
     1998-239725 [21]
                      WPIX
ΑN
DNC C1998-074761
TΤ
     New culture medium - includes, e.g.
     L-buthionine-(S,R)-sulphoximine, is useful in assessment of intracellular
     cysteine and glutathione concentrations.
DĊ
     B04 D16
IN
     CRAWFORD, J F
PA
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     128650 A IL 1997-128650 19970903
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     on WO 9810092; JP 2001500011 W Based on WO 9810092; KR 2000068398 A Based
     on WO 9810092; IL 128650 A Based on WO 9810092
PRAI US 1996-25373P
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IC
          C12Q001-02
     ICS
          C12N005-02; C12N005-06
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WO

AΒ

9810092 A UPAB: 19980528

A cell culture medium, which is useful for (i) determining levels of intracellular function of glutathione in lymphocytes and (ii) performing biochemical analysis of the antioxidant function of the lymphocytes, comprising a buffered, serum-free medium (pH 6.8-7.6) comprising: (a) a carbohydrate (which is glucose or a compound capable of producing this in the lymphocytes); (b) a biologically usable form of pantothenic acid, choline or a biologically usable form of a substance capable of producing choline in the lymphocytes; (c) inorganic ions comprising chloride, phosphate, calcium, magnesium, potassium, sodium and iron in a biologically utilisable form; (d) L-buthionine-(S,R)-sulphoximine (I); (e) deionised water, and (f) a mitogen in an amount effective to stimulate the lymphocytes. Also claimed is a cell culture medium, which is useful for (i) determining levels of intracellular function of **cysteine** and (ii) performing biochemical analysis of the antioxidant function of human lymphocytes, having identical composition to the medium above, but instead of (I) containing cumene hydroperoxide.

The media comprises a 5-500 mu M concentration of (I) or a 50-500 mu M concentration of (II). The media may be supplemented with amino acids and/or vitamins. The amino acids are selected from L-arginine, L-cysteine, L-glutamine, glycine, L-histidine, L-isoleucine, L-leucine, L-lysine, L-methionine, L-phenylalanine, L-serine, L-threonine, L-tryptophan, L-tyrosine and L-valine. The vitamins are selected from biotin, folinic acid, nicotinamide, nicotinic acid, riboflavin, thiamine, vitamin B6 and vitamin B12. Processes in which the media are used typically comprise: (a) inoculating the medium with lymphocytes from an individual; (b) incubating the inoculated medium, and (c) comparing the response of the lymphocytes with an average response of lymphocytes from a control group of individuals.

USE - The media may be used in processes for measuring levels of intracellular function of **cysteine** and glutathione, so as to provide a measurement of an individual's ability to prevent degenerative disease and deal with oxidative stress, and to allow therapeutic measures to be taken to improve an individual's antioxidant profile. It is widely accepted that certain conditions (e.g. ageing, arthritis, cancer, atherosclerosis, myocardial infarction, stroke, viral infection, pulmonary conditions, bowel diseases and neurodegenerative disease) can develop due to the presence of reactive oxygen molecules. Dwg.0/0

FS CPI

FA AB; DCN

MC CPI: B04-F04; B05-A01A; B05-A01B; B05-A03A; B05-C07; B07-A02B; B10-A08; B10-A22; B10-B02D; B10-C04E; B12-K04; D05-H01; D05-H09

L127 ANSWER 2 OF 2 WPIX (C) 2003 THOMSON DERWENT

AN 1998-063159 [06] WPIX

DNC C1998-022167

TI Cell culture medium for analysing antioxidant function in human lymphocytes - comprises buffered solution comprising nutrients, cumene hydroperoxide and mitogen, useful in, e.g. assessing intracellular deficiencies of vitamin(s).

DC B04 D16

IN BUCCI, L; CRAWFORD, J F; CRAWFORD, F

PA (RERE-N) RES DEV FOUND

CYC 75

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N 20 DO AO 11 A1 M 10 10 DE 35 DE 35 DE 17 DA 17

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    19970618; ZA 9705359 A ZA 1997-5359 19970618; EP 925370 A1 EP 1997-930001
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         C12N005-06; C12Q001-08
         A01N001-02; C12N001-38; C12Q001-02; G01N001-00; G01N033-50
    ICS
AΒ
         9748821 A UPAB: 19980209
    Cell culture medium (CCM) for biochemical analysis of antioxidant function
     in human lymphocytes comprises buffered, serum-free solution of pH 6.8-7.6
    comprising: (a) glucose or its in vivo precursor; (b) an utilisable form
    of pantothenic acid; (c) choline or its in
    vivo precursor; (d) inorganic ions selected from chloride, phosphate,
    calcium, magnesium, potassium, sodium and iron; (e) cumene
    hydroperoxide; (f) deionised water, and (g) mitogen to stimulate
    the lymphocytes being analysed. Also claimed are: (1) a method of
    determining abnormal quantitative nutritional requirements for specific
    required nutrients in an individual comprising: (a) inoculating the CCM
    with lymphocytes from the individual, where the CCM comprises limiting
    concentrations of the nutrient being tested; (b) incubating the inoculated
    CCM, and (c) comparing the response of the lymphocytes with an average
    response of the lymphocytes with an average response of lymphocytes from
    the control group, and (2) a method of identifying nutritional factors or
    biochemical intermediates or their products and other blood components
    including drugs in an individual sensitive to such detrimental effects
    comprising: (a) (1)(a), but where the CCM comprises at least 1 of the
    above mentioned components; (b) (1)(b), and (c) (1)(c), but where the
    response is compared with that in the same medium with a source of the
    substance suspected to affect the detrimental effect of the nutrient,
    biochemical intermediate or its product or other blood component including
    the drug being tested.
         USE - The methods and the CCM are used to biochemically analyse the
    cellular function of antioxidants (claimed). They also may be used to how
    well nutrient systems are working, to assess intracellular vitamin
    deficiencies that limit mitogenic responses, to determine abnormal
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nutrient requirements and to identify agents that can overcome the adverse action of blood components including drugs. Optimisation of vitamin and other nutrient levels in cells may, e.g. prevent heart diseases and some forms of cancer, stimulate the immune system, slow aging processes and

have a beneficial effect in disorders such as alcoholism, arthritis, diabetes, human immunodeficiency virus/acquired immune deficiency syndrome, macular degeneration and osteoporosis.

ADVANTAGE - The method may detect nutritional deficiencies before these become clinically manifested. Unlike known static methods, this process determines activity within cells. Lymphocytes are easy to collect and stimulate, they reflect long-term nutrient status and also possess metabolic pathways common to other cells and provide patient-specific information.

Dwg.0/1

FS CPI

FA AB; DCN

MC CPI: B03-D; B06-D09; B06-F03; B11-C08E; B12-K04A; B14-F01; B14-H01; D05-H01; D05-H08; D05-H09

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FILE LAST UPDATED: 25 APR 2003 <20030425/UP>
PATENTS CITATION INDEX, COVERS 1973 TO DATE

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L128 ANSWER 1 OF 1 DPCI (C) 2003 THOMSON DERWENT

AN 1998-239725 [21] DPCI

DNC C1998-074761

TI New culture medium - includes, e.g. L-buthionine-(S,R)-sulphoximine, is useful in assessment of intracellular cysteine and glutathione concentrations.

DC B04 D16

IN CRAWFORD, J F

PA (RERE-N) RES DEV FOUND

CYC 78

PI WO 9810092 A1 19980312 (199821)\* EN 36p C12Q001-02 <-RW: AT BE CH DE DK EA ES FI FR GB GH GR IE IT KE LS LU MC MW NL OA PT
SD SE SZ UG ZW

W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GE HU IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK TJ TM TR TT UA UG UZ VN

AU 9742464 A 19980326 (199832) C12Q001-02 ZA 9707892 A 19990428 (199922) 35p A61K000-00 A1 19990728 (199934) EN C12Q001-02 EP 931163 R: AT BE CH DE DK ES FI FR GB GR IE IT LI NL PT SE B 20000420 (200029) · C12Q001-02 AU 718816 NZ 334327 A 20000623 (200038) C12Q001-02 CN 1268977 A 20001004 (200067) C12Q001-02 JP 2001500011 W 20010109 (200107) 33p C12Q001-02 KR 2000068398 A 20001125 (200130) C12Q001-02

KR 2000068398 A 20001125 (200130) C12Q001-02 US 2002068270 A1 20020606 (200241) C12Q001-00 IL 128650 A 20021201 (200282) C12N005-00

ADT WO 9810092 A1 WO 1997-US15451 19970903; AU 9742464 A AU 1997-42464 19970903; ZA 9707892 A ZA 1997-7892 19970903; EP 931163 A1 EP 1997-940761 19970903, WO 1997-US15451 19970903; AU 718816 B AU 1997-42464 19970903; NZ 334327 A NZ 1997-334327 19970903, WO 1997-US15451 19970903; CN 1268977 A CN 1997-197585 19970903; JP 2001500011 W WO 1997-US15451 19970903, JP 1998-512821 19970903; KR 2000068398 A WO 1997-US15451 19970903, KR 1999-701715 19990302; US 2002068270 A1 Provisional US 1996-25373P 19960903, Div ex US 1997-922279 19970903, US 2001-17625 20011213; IL 128650 A IL 1997-128650 19970903

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PRAI US 1996-25373P 19960903; US 1997-922279 19970903; US 2001-17625 20011213

IC ICM A61K000-00; C12N005-00; C12Q001-00; C12Q001-02 ICS C12N005-02; C12N005-06

FS CPI

# CTCS CITATION COUNTERS

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# Cited by Examiner

CITING PATENT	CA:	CITED PATENT ACCNO
WO 9810092		US 4927762 A 1990-185660/24
	PA:	(CELL-N) CELL ENTERPRISES
	IN:	DARFLER, F J
	Y	US 5171885 A 1993-008659/01
	PA:	(CORR) CORNELL RES FOUND INC
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	Y	US 5290571 A 1994-074311/09
	PA:	(IMMU-N) IMMUNOTEC RES CORP LTD
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	PA:	(KURB) KURASHIKI BOSEKI KK; (KURB) KURABO IND LTD
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L129 ANSWER 1 OF 1 DPCI (C) 2003 THOMSON DERWENT
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AN 1998-063159 [06] DPCI

DNC C1998-022167

TI Cell culture medium for analysing antioxidant function in human lymphocytes - comprises buffered solution comprising nutrients, cumene hydroperoxide and mitogen, useful in, e.g. assessing intracellular deficiencies of vitamin(s).

DC B04 D16

IN BUCCI, L; CRAWFORD, J F; CRAWFORD, F

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                                                    C12Q001-08
    EP 925370 B1 20021218 (200301) EN
                                                    C12Q001-08
        R: AT BE CH DE DK ES FI FR GB GR IE IT LI NL PT SE
     DE 69718017 E 20030130 (200317)
                                                    C12Q001-08
    WO 9748821 A1 WO 1997-US10328 19970618; AU 9733934 A AU 1997-33934
ADT
    19970618; ZA 9705359 A ZA 1997-5359 19970618; EP 925370 A1 EP 1997-930001
    19970618, WO 1997-US10328 19970618; CN 1222940 A CN 1997-195713 19970618;
    US 5985665 A US 1996-665941 19960619; NZ 333231 A NZ 1997-333231 19970618,
    WO 1997-US10328 19970618; AU 720703 B AU 1997-33934 19970618; JP
    2000514287 W WO 1997-US10328 19970618, JP 1998-503167 19970618; IL 127576
    A IL 1997-127576 19970618; KR 2000016773 A WO 1997-US10328 19970618, KR
    1998-710382 19981218; EP 925370 B1 EP 1997-930001 19970618, WO
    1997-US10328 19970618; DE 69718017 E DE 1997-618017 19970618, EP
    1997-930001 19970618, WO 1997-US10328 19970618
FDT AU 9733934 A Based on WO 9748821; EP 925370 Al Based on WO 9748821; NZ
     333231 A Based on WO 9748821; AU 720703 B Previous Publ. AU 9733934, Based
    on WO 9748821; JP 2000514287 W Based on WO 9748821; KR 2000016773 A Based
     on WO 9748821; EP 925370 Bl Based on WO 9748821; DE 69718017 E Based on EP
     925370, Based on WO 9748821
PRAI US 1996-665941
                     19960619
    ICM A61K000-00; C12N005-00; C12N005-02; C12N005-06; C12Q001-08
    ICS
         A01N001-02; C12N001-38; C12Q001-02; G01N001-00; G01N033-50
FS
    CPI
EXF EXAMINER'S FIELD OF SEARCH
                                UPE: 20000113
   US 5985665
                  A 19991116
     435/014; 435/002; 435/029; 435/372; 435/375; 435/387; 435/004;
     435/040.500; 435/404; 435/405; 435/406
CTCS CITATION COUNTERS
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Page 54 gitomer - 10 / 017625

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UPD: 20000113 CDP CITED PATENTS

Cited by Examiner

CITING PATENT CAT CITED PATENT ACCNO \_\_\_\_\_

WO 9748821 A 'No Citations US 5985665 A US 4499064 A 1983-846286/51

PA: (RERE-N) RES DEV FOUND; (CLAY-N) CLAYTON FOUND RES

IN: SHIVE, W

US 4927762 A 1990-185660/24

PA: (CELL-N.) CELL ENTERPRISES

IN: DARFLER, F J

REN LITERATURE CITATIONS UPR: 19980603

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Citations by Examiner \_\_\_\_\_\_

CITING PATENT CAT CITED LITERATURE

\_\_\_\_\_\_ WO 9748821 A ARCH. BIOCHEM. BIOPHYS., 1976, Vol. 175, NORDBLOM

et al., "Studies on Hydroperoxide-Dependent Substrate Hydroxylation by Purified Liver

Microsomal Cytochrome P-450", pages 524-533. ANALYTICAL BIOCHEMISTRY, 1992, Vol. 202, JIANG et WO 9748821 A

al., "Ferrous Ion Oxidation in the Presence of

Xylenol Orange for Detection of Lipid

Hydroperoxide in Low Density Lipoprotein", pages

384-389.

CGP CITING PATENTS UPG: 20010409

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Cited by Examiner

CITED PATENT CAT CITING PATENT ACCNO

US 5985665 A US 6165797 A 2001-158205/12

PA: (BIOD-N) BIO DEFENSE NUTRITIONALS INC

IN: HALSTEAD, B W

=> fil wpix

FILE 'WPIX' ENTERED AT 17:38:01 ON 26 APR 2003

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16 APR 2003 <20030416/UP> FILE LAST UPDATED:

MOST RECENT DERWENT UPDATE: 200325 <200325/DW>

DERWENT WORLD PATENTS INDEX SUBSCRIBER FILE, COVERS 1963 TO DATE

Due to data production problems the WPI file had to be reset to update 200324.

SDIs for update 24 will be rerun.

The previous SDI run for 24 has been credited.

Also answer sets created after April 10 may at least temporarily be affected and hence partially invalid.

>>> NEW WEEKLY SDI FREQUENCY AVAILABLE --> see NEWS <<<

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>>> SLART (Simultaneous Left and Right Truncation) is now
    available in the /ABEX field. An additional search field
    /BIX is also provided which comprises both /BI and /ABEX <<<
>>> PATENT IMAGES AVAILABLE FOR PRINT AND DISPLAY <<<
>>> FOR DETAILS OF THE PATENTS COVERED IN CURRENT UPDATES,
    SEE http://www.derwent.com/dwpi/updates/dwpicov/index.html <<<
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    http://www.derwent.com/userguides/dwpi guide.html <<<
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                           (C) 2003 THOMSON DERWENT
L132 ANSWER 1 OF 7 WPIX
     2001-158205 [16]
                        WPIX
DNN
    N2001-115196
                        DNC C2001-046850
ΤI
    Detection of malondialdehyde as indicator of oxidative stress, in urine,
    using home test kit including basic fuchsin.
DC
    B04 S03
IN
    HALSTEAD, B W
PΑ
     (BIOD-N) BIO DEFENSE NUTRITIONALS INC
CYC
                                                     G01N021-78
PΙ
    US 6165797
                 A 20001226 (200116)*
                                               3p
                                                                     <--
ADT
    US 6165797 A US 1999-253223 19990219
PRAI US 1999-253223
                      19990219
IC
     ICM G01N021-78
     ICS
         G01N033-493
AΒ
          6165797 A UPAB: 20010323
    NOVELTY - Malondialdehyde (I) is detected in urine using a reagent
     comprising acetic acid, sodium metabisulfite, phosphoric acid, basic
     fuchsin and deionized water.
          DETAILED DESCRIPTION - Using a test kit, which comprises a reagent,
     to detect malondialdehyde (I) in urine comprises:
          (1) combining the reagent with the urine sample to produce a colored
    product and
          (2) comparing the color of the product with a reference chart to
    determine the amount of (I) in the sample.
         The reagent comprises:
          (i) 90-110 parts of 20% acetic acid;
          (ii) 13.5-16.5 parts of ingredient A; and
          (iii) 4.5-5.5 parts of ingredient B.
          Ingredient A comprises 18-22 g of sodium metabisulfite, 9-11 ml of
     concentrated phosphoric acid and 450-550 ml of deionized water.
          Ingredient B comprises 0.45-0.55 g of basic fuchsin and 90-110 ml of
     ingredient A.
          USE - The process is useful for detection of oxidative stress.
          ADVANTAGE - The fuchsin based colorimetric test is rapid and easy to
     carry out, and can be performed using a home test kit with a small
     quantity of urine. Retesting after a suitable period of time can assess
     the adequacy of antioxidant therapy. A color change can typically be
     observed when the concentration of (I) in urine is greater than 2 ppm.
     Dwg.0/0
FS
     CPI EPI
FΑ
    AB; DCN
```

CPI: B04-B04B1; B05-A01B; B05-B02A3; B10-A01; B10-D01; B11-C07B1; B12-K04A

MC

EPI: S03-E04E; S03-E14H

TECH UPTX: 20010323 TECHNOLOGY FOCUS - BIOLOGY - Preferred process: The reagent especially comprises 100 parts of 20% acetic acid, 15 parts of ingredient A and 5  $\,$ parts of ingredient B. The proportions in ingredient A are especially 20 g of sodium metabisulfite, 10 ml of phosphoric acid and 500 ml of deionized water. The proportions in ingredient B are especially 0.5 q of basic fuchsin to 100 ml of ingredient A. After the reagent is combined with the urine sample, the mixture is allowed to stand for at least 2 minutes before the color is compared with the reference chart. The test reagent is sealed in an ampoule, then the test sample is injected into the ampule using a plastic bulb. L132 ANSWER 2 OF 7 WPIX (C) 2003 THOMSON DERWENT 1995-052070 [07] WPIX AN C1995-023911 DNC ΤI Serum-depleted or serum-free culture medium for long-term cell growth contg. albumin, transferrin, nucleoside(s), growth factor, extracellular matrix material, pyruvate, cholesterol, standard medium etc.. DC B04 D16 IN PONTING, I L O PA (AMGE-N) AMGEN INC CYC 49 PΙ WO 9500632 A1 19950105 (199507) \* EN 60p C12N005-06 RW: AT BE CH DE DK ES FR GB GR IE IT LU MC NL OA PT SE W: AT AU BB BG BR BY CA CH CN CZ DE DK ES FI GB HU JP KP KR KZ LK LV MG MN MW NL NO NZ PL PT RO RU SD SE SK UA UZ VN US 5405772 A 19950411 (199520) 22p C12N005-00 <--AU 9471124 Α 19950117 (199521) C12N005-06 EP 703978 A1 19960403 (199618) EN C12N005-06 R: AT BE CH DE DK ES FR GB GR IE IT LI LU MC NL PT SE JP 08508891 W 19960924 (199704) 52p C12N005-06 В AU 678836 19970612 (199732) C12N005-06 MX 186150 B 19970926 (199850) C12N005-000 JP 2866742 B2 19990308 (199915) 25p C12N005-06 EP 703978 B1 19990818 (199937) EN C12N005-06 R: AT BE CH DE DK ES FR GB GR IE IT LI LU MC NL PT SE DE 69420138 E 19990923 (199945) C12N005-06 ES 2135589 T3 19991101 (199953) C12N005-06 WO 9500632 A1 WO 1994-US6893 19940617; US 5405772 A US 1993-79719 ADT 19930618; AU 9471124 A AU 1994-71124 19940617; EP 703978 A1 EP 1994-920264 19940617, WO 1994-US6893 19940617; JP 08508891 W WO 1994-US6893 19940617, JP 1995-502993 19940617; AU 678836 B AU 1994-71124 19940617; MX 186150 B MX 1994-4604 19940617; JP 2866742 B2 WO 1994-US6893 19940617, JP 1995-502993 19940617; EP 703978 B1 EP 1994-920264 19940617, WO 1994-US6893 19940617; DE 69420138 E DE 1994-620138 19940617, EP 1994-920264 19940617, WO 1994-US6893 19940617; ES 2135589 T3 EP 1994-920264 19940617 FDT AU 9471124 A Based on WO 9500632; EP 703978 A1 Based on WO 9500632; JP 08508891 W Based on WO 9500632; AU 678836 B Previous Publ. AU 9471124, Based on WO 9500632; JP 2866742 B2 Previous Publ. JP 08508891, Based on WO 9500632; EP 703978 B1 Based on WO 9500632; DE 69420138 E Based on EP 703978, Based on WO 9500632; ES 2135589 T3 Based on EP 703978 PRAI US 1993-79719 19930618 REP 1.Jnl.Ref; WO 9218615; WO 9309220 ICM C12N005-00; C12N005-000; C12N005-06 IC C12N005-002; C12N005-006; C12N005-08 ICS C12N005-06, C12R001:91 ICI 9500632 A UPAB: 19960625

A serum-depleted or serum-free medium for the long-term proliferation and development of cells comprises (a) a standard culture medium, (b) serum albumin, (c) transferrin, (d) a source of lipids and fatty acids, (e) cholesterol, (f) a reducing agent, (g) pyruvate, (h) nucleosides for synthesis of DNA and RNA, (i) at least one growth factor that stimulates the proliferation and development of stromal cells, tissue cells or organ

cells, and (j) at least one extracellular matrix (ECM) material. USE - The medium can be used for both short and long-term maintenance of proliferation and development of cells, including fibroblasts, glial cells, neuronal cells, adipocytes, myoblasts, epithelial cells, hepatocytes, osteoclasts, heart muscle cells and lymphopoietic cells, and partic. haematopoietic cells. The medium can be used to stimulate the proliferation and/or development of early progenitor cells for bone marrow transplants and/or gene transfer into these cells for gene therapy for treating immunological or haematological disorders, e.g. severe combined immunodeficiency, adenosine deaminase deficiency and AIDS. The medium can also be used to determine the function of a novel gene by adding an anti-sense oligomer to inhibit expression. ADVANTAGE - The chemically defined medium can provide growth of primal cells as well as immortalised cell lines and development for up to several months. The medium allows the growth of hemopoietic cells and the stromal cells that support them. Dwg.0/7 CPI AB; DCN CPI: B01-D02; B04-B01B; B04-B04D4; D05-H01 ABEQ US 5405772 A UPAB: 19950530 Medium for long-term proliferation and development of cells comprises (a) 0.8-1.09 times standard culture medium; (b) 3-50 mg per ml serum albumin; (c) 25-1000 micro-g/ml transferrin; (d) 5-100 micro-g/ml lipids and fatty acids; (e) 3-30 micro-g/mol cholesterol; (f) 30-300 microM reducing agent; (g) 30-500 micro-g/ml pyruvate; (h) 5-30 micro-g/ml nucleosides; (i) growth factor; and (j) extracellular matrix material(s). Cpd. (i) comprises 5-200 ng per mol. epidermal growth factor, 0.5-40 ng per ml fibroblast growth factor, 2-200 ng/ml platelet-derived growth factor, and/or 2-100 micro-g/ml insulin. Cpd. (j) comprises 2-100 micro-g/cm2 collagen IV and/or 0.5-100 micro-g/cm2 fibronectin. USE - Used for culturing adipocytes, macrophages, endothelial cells, fibroblasts and haematopoietic progenitor cells. Dwg.0/8 L132 ANSWER 3 OF 7 WPIX (C) 2003 THOMSON DERWENT 1994-074311 [09] WPIX 1989-317523 [44]; 1990-194723 [26]; 1991-231513 [32]; 1993-351356 [44]; 1994-008103 [02]; 1995-365221 [43] C1994-033793 Whey protein compsn. useful to improve humoral immune response - contq. undenatured whey protein concentrate opt. with vitamins B1 and B2. BOUNOUS, G; GOLD, P; KONGSHAVN, P A L (IMMU-N) IMMUNOTEC RES CORP LTD US 5290571 A 19940301 (199409)\* 19p A61K035-20 US 5290571 A Cont of US 1988-188271 19880429, CIP of US 1988-289971 19881223, US 1989-417246 19891004 PRAI US 1989-417246 19891004; US 1988-188271 19880429; US 1988-289971 19881223 ICM A61K035-20 5290571 A UPAB: 19951204 US Whey protein compsn. (I) comprises undentured whey protein concentrate obtd. from raw borine, goat or sheep milk and contq. oil and heat labile whey protein present in the raw milk, in an amt. of 18-28g of whey protein per 100g of compsn., vitamin B1 and in amt. of at least 1.5 (pref. 1.5-2.0) mg per 100g compsn. and Vitamin B2 in an amt. of at least 1.5 (pref. 1.5-2.0)mg per 100g compsn.. Also claimed is a method of improving the immune response in mammals

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ADT

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AB

as measured by sheep red blood cell injection by oral admin. of (I), the vitamins B1 and B2 being admin. in amts. in excess of minimum daily requirements. Embodiments also claimed include admin. of the compsn.

comprising vitamin B2 and an amt. in excess of minimum daily requirements and the undenatured whey protein concentrate in an amt. sufficient to satisfy the daily requirements of protein of the mammal; and admin. of the whey protein concentrate having immuno-enhancing properties which are heat labile, insensitive to pancreatic digestion and dependent upon the undenatured state.

USE - Oral admin. of (I) increases the concn. level of glutathione in the organs of mammals and enhances resistance to bacterial infection, partic. pneumococcal infection and slow growing carcinoma such as colon carcinoma.

In an example, C3H/HeJ mice fed a diet contg. 20g undenatured whey protein (U-Lacp)/100g diet showed improved survival after i.v. infection with Streptococcus pneumanial type 3 and compared to similarly infected mice fed a 20g C/diet of similar nutritional efficiency.

Dwg.0/12 Dwg.0/12

Dwg.u/

FS CPI

FA AB; DCN

MC CPI: B03-B; B03-C; B04-N02; B14-G01; D03-B02

L132 ANSWER 4 OF 7 WPIX (C) 2003 THOMSON DERWENT

AN 1993-008659 [01] WPIX

CR 1993-302719 [38]; 1994-091597 [11]

DNC C1993-004029

TI New 2-homo cysteine-5-sulphoximine and R-diastereoisomer and homologues - causes depletion of glutathione esp. in heart and liver.

DC B05

IN GRIFFITH, O W

PA (CORR) CORNELL RES FOUND INC

CYC 1

PI US 5171885 A 19921215 (199301)\* 6p C07C053-128 <--

ADT US 5171885 A Cont of US 1989-359886 19890601, US 1991-715898 19910619

PRAI US 1989-359886 19890601; US 1991-715898 19910619

IC ICM C07C053-128

AB US 5171885 A UPAB: 19940428

Pure L-(S-(S-6C alkyl)) homocysteine-S-sulphoximes and their acid-addition salts are new.

L-Buthionine-S-sulphoxime and the diastereoisomeric L-buthionine-R-sulphoxime, in pure form, and their acid-addition salts, are specifically claimed. These cpds. have the formulae (I) and (II) respectively.

USE - (I) has use, e.g. in causing the depletion of glutathione, a major protectant molecule in tumours and certain parasites. (II) has use as a cpd. decreasing further uptake of (I) by heart and liver. Since decreased uptake of (I) is associated with smaller decreases in glutathione content, heart and liver would be partially protected from the adverse effects of glutathione depletion. The pure diastereoisomer can be administered alone or with a carrier either orally or parenterally, e.g. at daily dosages in the range 0.1-30 mM/kg.

Dwg.0/0

13

Dwg.0/0

FS CPI

FA AB; GI; DCN

MC CPI: B10-A01

L132 ANSWER 5 OF 7 WPIX (C) 2003 THOMSON DERWENT

AN 1992-293906 [36] WPIX

DNC C1992-130644

TI Serum-free media for culturing animal cells - contg. specified amt. of methionine.

DC B04 D16

IN ARAKAWA, H; NISHINO, T; SAKAI, C; TORISHIMA, H; YAMAMOTO, R

```
(KURB) KURASHIKI BOSEKI KK; (KURB) KURABO IND LTD
PΑ
CYC
    7
PΙ
     EP 501435
                   A1 19920902 (199236) * EN
                                              20p
                                                     C12N005-00
         R: DE FR GB IT NL
     JP 04271779
                  A 19920928 (199245)
                                              15p
                                                     C12N005-06
     US 5326699
                   A 19940705 (199426)
                                               6p
                                                     C12N005-00
                                                                      <--
     JP 07089908
                   B2 19951004 (199544)
                                              11p
                                                     C12N005-06
     EP 501435
                   B1 19981028 (199847)
                                         ΕN
                                                     C12N005-00
         R: DE FR GB IT NL
     DE 69227392
                  E 19981203 (199903)
                                                     C12N005-00
ADT
    EP 501435 A1 EP 1992-103224 19920226; JP 04271779 A JP 1991-34150
     19910228; US 5326699 A Cont of US 1992-842980 19920228, US 1993-120235
     19930914; JP 07089908 B2 JP 1991-34150 19910228; EP 501435 B1 EP
     1992-103224 19920226; DE 69227392 E DE 1992-627392 19920226, EP
     1992-103224 19920226
FDT JP 07089908 B2 Based on JP 04271779; DE 69227392 E Based on EP 501435
PRAI JP 1991-34150
                      19910228
    5.Jnl.Ref; EP 354129; FR 1283157; US 4767704; WO 9007007
IC
         C12N005-06
AΒ
           501435 A UPAB: 19931112
     EΡ
     Serum-free media for culturing animal cells contain 8-14 mg/l of
    methionine (I).
          The compsns. contain 5-11.5 mg/l threonine, 7-11 mg/l tyrosine,
     9-14.5 mg/l phenylalanine, 400-500 mg/l glutamine, 7-9.5 mg/l aspartic
     acid, 19-28 mg/l lysine.HCl, 33-40 mg/l serine and 0-0.1 mM CaCl2 (N.B.,
     'mg/l' is given as 'mg/ml' in the claims).
          USE/ADVANTAGE - The media are esp. useful for culturing epithelial
     cells, e.g., epidermal keratinocytes and corneal epithelial cells. The
    media provide better growth than media contg. different amts. of (I)
     Dwq.0/0
FS
    CPI
FA
    AB; DCN
     CPI: B04-B04A3; B10-B02D; B11-A; D05-H01; D05-H08
MC.
          5326699 A UPAB: 19940817
     Serum free medium for culturing animal epithelial cells comprises 9-14
    mg/l methionine, 5-11.5 mg/l phenylalanine, 400-550 mg/l glutamine, 7-9.5
    mg/l aspartic acid, 19-28 mg/l lysine HCl, 33-40 mg/l serine and up to 0.1
    mM Ca in the form CaCl2, and one or more conventional ingredients selected
     from glucose, vitamins, minerals, growth factors, HEPES and amino acids
     other than those above pref. ala, arg, asp, cys, glutamic acid, gly, his,
     ile, leu, pro, try, val and their alkaline or and salts.
          USE - For culturing animal cells pref. epithelial cells.
     Dwq.0/0
L132 ANSWER 6 OF 7 WPIX
                           (C) 2003 THOMSON DERWENT
     1990-185660 [24]
                        WPIX
AN
DNC
    C1990-080475
ΤI
     Chemically defined, serum-free medium for cells - contg. cpd. having one
     free thiol gp. to remove toxicity due to oxidising agents present in basal
     media.
DC
     B04 B05 D16
ΙN
     DARFLER, F J
PA
     (CELL-N) CELL ENTERPRISES
CYC
    1
PΙ
                  A 19900522 (199024)*
                                                                      <--
     US 4927762
    US 4927762 A US 1988-281974 19881130
ADT
PRAI US 1986-846716
                      19860401; US 1988-281974
                                                 19881130
TC
     C12N001-38; C12N005-00
          4927762 A UPAB: 19930928
     A chemically defined, serum-free medium for the maintenance or growth of
     immortal or immortalised cells is claimed which comprises a cell growth
     promoting amt. of a cpd. (I) having one free thiol gp. and which is
     capable of supporting the growth of immortal or immortalised cells at seed
```

densities less than x 10 power (5) cells/ml. Pref. (I) is of formula (Ia) SH-CR1R2-CR3R4R5 (Ia) (R1, R4 = H, NH2 or NH (CO) CH3; R5 = COO-, SO3-, or CH2SO3- esp. thiolactate. The medium may also contain insulin or transferrin. Also claimed is a supplemental cell culture medium consisting of transferrin, insulin, ethanolamine, a selenium salt, an unsatd. fatty acid and a cell-growth promoting amt. of a cpd. (II) having one free thiol gp. where the medium, when added to a basal cell culture medium, supports the growth of hybridoma cells at seed densities of less than 10 5 cells/ml. (II) may be e.g. N-acetylcysteine, D- or L-pencillamine, 2-mercaptothanesulphonic acid (MENSA' or mercaptopropionic acid (MPA). ADVANTAGE - The cpds. are protective agents that remove toxicity due to oxidising agents ordinarily present in basal tissue culture media. They support the long-term growth of lymphoid cells, including hybridomas, even at low seed densities. @ 0/0 CPI AB; DCN CPI: B04-B02D2; B04-B04A6; B05-B02C; B10-A09B; B10-B02D; B10-B04B; B10-C04D; B10-C04E; B11-A; D05-H01 L132 ANSWER 7 OF 7 WPIX (C) 2003 THOMSON DERWENT 1983-846286 [51] WPIX C1983-123493 Culture medium contg. nutrients, metals, vitamin(s) etc. - useful in lymphocyte assays for nutritional status in subject. B04 D16 SHIVE, W (RERE-N) RES DEV FOUND; (CLAY-N) CLAYTON FOUND RES 19831221 (198351) \* EN 35p R: AT BE CH DE FR IT LI LU NL SE AU 8315364 A 19831208 (198405) GB 2124366 19840215 (198407) NO 8301985 19831227 (198407) DK 8302543 19840123 (198411) FI 8302005 Α 19840131 (198411) ZA 8304018 Α 19840216 (198426) PT 76808 Α 19841018 (198447) ES 8405953 A 19841001 (198449) <--A 19850212 (198509) US 4499064 A 19860128 (198609) CA 1199883 в 19861112 (198646) GB 2124366 EP 96560 B 19870506 (198718) EN R: AT BE CH DE FR IT LI LU NL SE DE 3371362 G 19870611 (198724) IL 68873 Α 19870331 (198724) KR 8900729 В 19890330 (198941) DK 172418 В 19980602 (199828) C12N005-02 EP 96560 A EP 1983-303225 19830603; GB 2124366 A GB 1983-15287 19830603; ZA 8304018 A ZA 1983-4018 19830603; US 4499064 A US 1983-492308 19830506; DK 172418 B DK 1983-2543 19830603 DK 172418 B Previous Publ. DK 8302543 19820603; US 1983-492308 PRAI US 1982-383822 19820603; US 1982-384822 19830506 EP 66284; US 3128228 A61B010-00; C12N005-00; C12Q001-04; C12Q003-00; G01N001-00; G01N033-50

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REP

ICS

C12N005-02

G01N001-00; G01N033-50

DNC

AB EΡ 96560 A UPAB: 19930925 Culture medium for quantitative lymphocyte assay of the nutritional status of an individual comprises a buffered serum-free soln., including (1)

A61B010-00; C12N005-00; C12Q001-02; C12Q001-04; C12Q003-00;

glucose or a substance capable of producing it as a metabolic prod.; (2) the amino acids arginine, cysteine, glutamine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, serine, threonine, tryptophan, tyrosine and valine(all in L-form) and glycine, these acids being present as a gp. each in an amount up to the amount in a normal blood range; (3) pantothenate opt. with biotin, folinic acid, nicotinamide or nicotinic acid, riboflavin, thiamine, vitamin B6 and/or vitamin B12; (4) Cl, SO4, PO4, Ca, Mg, K, Na and ferrous ions; (5) adenine and choline or their precursors, inositol, and pyruvate or another effective metabolite; (6) a nitrogen to stimulate lymphocytes being assayed; and (7) deionised water. The buffer is at pH 6.8-7.6.

Lymphocytes from small blood samples are readily available on a routine basis and are metabolically inactive until activated by a mitogen, so that they carry information about past nutritional status and have little day-to-day variation in nutritional responses.

0/0

FS CPI

FA AB

MC CPI: B04-A07F; B04-B04A; B04-B04D; B11-C07; B12-K04; D05-H01

ABEQ EP 96560 B UPAB: 19930925

A cell culture medium for culturing lymphocytes comprising a buffered, serum-free solution in deionised water, of the following components:- (a) glucose or a carbohydrate which is capable of being converted metabolically to glucose; (b) choline or a precursor of choline; (c) an inorganic ion supplement comprising chloride, phosphate, calcium, magnersium, potassium, sodium and iron ions; (d) a mitogen to stimulate the metabolic activity of lymphocytes; and (e) an amino acid supplement comprising L-argine, L-cysteine, L-glutamine, glycine, L-histidine, L-isoleucine, L-leucine, L-lysine, L-methionine, L-phenylalanine, L-serine, L-threonine, L-tryptophan, L-tyrosine and L-valine, and/or (f) a vitamin supplement comprising one or more vitamins selected from pantothenic acid, biotin, folinic acid, nicotinamide or nicotinic acid, riboflavin, thiamin, vitamin B and vitamin B12; said buffered, serum-free solution having a pH in the range of from 6.8 to 7.6. GB 2124366 B UPAB: 19930925

A cell culture medium for culturing lymphocytes comprising a buffered, serum-free solution in deionised water, of the following components:- a) glucose or a carbohydrate which is capable of being converted metabolically to glucose; b) choline or a precursor of choline; c) an inorganic ion supplement comprising chloride, sulphate, phosphate, calcium, magnesium, potassium, sodium and iron ions; d) a mitogen to stimulate the metabolic activity of lymphocytes; and e) an amino acid supplement comprising L-arginine, L-cysteine, L-glutamine, glycine, L-histidine, L-isoleucine, L-leucine, L-lysine, L-methionine, L-phenylalanine, L-serine, L-threonine, L-tryptophan, L-tyrosine and L-valine; and f) a vitamin supplement comprising one or more vitamins selected from pantothenic acid, biotin, folinic acid, nicotinamide or nicotinic acid, riboflavin, thiamin, vitamin B6 and vitamin B12; said buffered serum-free solution having a pH in the range of from 6.8 to 7.6. ABEQ US 4499064 A UPAB: 19930925

Cell culture medium comprises a buffered, serum-free soln. contg. (1) glucose or cpd. biologically capable of producing glucose in the cells, (2) biologically usable form of pantothenic acid, (3) choline or a substance producing it in the cells, (4) Cl, PO4, Ca, Mg, K, Na and Fe ions, (5) deionised water and (6) a mitogen to stimulate lymphocytes. The soln. has pH 6.8-7.6.

USE - For lymphocyte assay of nutritional and biochemical status of cells from human beings. Nutritional and biochemical deficiencies and inadequacies and imbalances of the lymphocytes can be determined when the medium is supplemented with an aminoacid or vitamin nutrient supplement. The nutrient being tested is omitted from or is present in limiting or inhibitory amts. in the nutrient supplement.

=> fil medline FILE 'MEDLINE' ENTERED AT 17:39:43 ON 26 APR 2003 FILE LAST UPDATED: 26 APR 2003 (20030426/UP). FILE COVERS 1958 TO DATE. On April 13, 2003, MEDLINE was reloaded. See HELP RLOAD for details.

MEDLINE thesauri in the /CN, /CT, and /MN fields incorporate the MeSH 2003 vocabulary. See http://www.nlm.nih.gov/mesh/changes2003.html for a description on changes.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d all 1133 L133 ANSWER 1 OF 1 MEDLINE ΑN 76267486 MEDLINE DN 76267486 PubMed ID: 8710 ΤI Studies on hydroperoxide-dependent substrate hydroxylation by purified liver microsomal cytochrome P-450. ΑU Nordblom G D; White R E; Coon M J SO ARCHIVES OF BIOCHEMISTRY AND BIOPHYSICS, (1976 Aug) 175 (2) 524-33. Journal code: 0372430. ISSN: 0003-9861. CY United States DT Journal; Article; (JOURNAL ARTICLE) English LA FS Priority Journals EM197610 Entered STN: 19900313 ED Last Updated on STN: 19970203 Entered Medline: 19761029 CTCheck Tags: Animal; Support, U.S. Gov't, Non-P.H.S.; Support, U.S. Gov't, P.H.S. \*Cytochrome P-450 Enzyme System: ME, metabolism Hydrogen-Ion Concentration Kinetics Microsomes, Liver: DE, drug effects \*Microsomes, Liver: EN, enzymology \*Mixed Function Oxygenases: ME, metabolism Oxidoreductases, N-Demethylating: ME, metabolism Oxygen \*Peroxides: PD, pharmacology Rabbits Structure-Activity Relationship ŔN 7782-44-7 (Oxygen); 9035-51-2 (Cytochrome P-450 Enzyme System) O (Peroxides); EC 1.- (Mixed Function Oxygenases); EC 1.5. (Oxidoreductases, N-Demethylating)

#### => d all 1134

- L134 ANSWER 1 OF 1 MEDLINE
- 92391672 MEDLINE ΑN
- DN 92391672 PubMed ID: 1519766
- TIFerrous ion oxidation in the presence of xylenol orange for detection of lipid hydroperoxide in low density lipoprotein.
- ΑU Jiang Z Y; Hunt J V; Wolff S P
- Department of Clinical Pharmacology, University College and Middlesex CS School of Medicine, London.
- SO ANALYTICAL BIOCHEMISTRY, (1992 May 1) 202 (2)

#### 384-9.

Journal code: 0370535. ISSN: 0003-2697.

- CY United States
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals
- EM 199210
- ED Entered STN: 19921023

Last Updated on STN: 19921023 Entered Medline: 19921007

- A simple and sensitive method for the direct measurement of lipid peroxides in lipoprotein and liposomes is described. The method is based on the principle of the rapid peroxide-mediated oxidation of Fe2+ to Fe3+ under acidic conditions. The latter, in the presence of xylenol orange, forms a Fe(3+)-xylenol orange complex which can be measured spectrophotometrically at 560 nm. Calibration with standard peroxides, such as hydrogen peroxide, linoleic hydroperoxide, t-butyl hydroperoxide, and cumene hydroperoxide gives a mean apparent extinction coefficient of  $4.52 \times 10(4)$  M-1 cm-1 consistent with a chain length of approximately 3 for ferrous ion oxidation by hydroperoxides. Endoperoxides are less reactive or unreactive in the assay. The assay has been validated in the study of lipid peroxidation of low density lipoprotein and phosphatidyl choline liposomes. By pretreatment with enzymes known to metabolize peroxides, we have shown that the assay measures lipid hydroperoxides specifically. Other methods for measuring peroxidation, such as the assessment of conjugated diene, thiobarbituric acid reactive substances and an iodometric assay have been compared with the ferrous oxidation-xylenol orange assay.
- CT Check Tags: Human; Support, Non-U.S. Gov't
  - \*Ferrous Compounds: AN, analysis
    - \*Fluorescent Dyes
    - \*Linoleic Acids: AN, analysis
    - \*Lipid Peroxides: AN, analysis
    - Lipoproteins, LDL: BL, blood \*Lipoproteins, LDL: CH, chemistry

Liposomes

Oxidation-Reduction

- \*Xylenes: CH, chemistry
- RN 1611-35-4 (xylenol orange); 25657-09-4 (linoleic acid hydroperoxide)
- CN 0 (Ferrous Compounds); 0 (Fluorescent Dyes); 0 (Linoleic Acids); 0 (Lipid Peroxides); 0 (Lipoproteins, LDL); 0 (Liposomes); 0 (Xylenes)

### => fil hcaplus

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FILE COVERS 1907 - 26 Apr 2003 VOL 138 ISS 18 FILE LAST UPDATED: 25 Apr 2003 (20030425/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

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=> d all
L135 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2003 ACS
     1976:474184 HCAPLUS
AN
     85:74184
DN
     Studies on hydroperoxide-dependent substrate hydroxylation by purified
TI
     liver microsomal cytochrome P-450
     Nordblom, Gerald D.; White, Ronald E.; Coon, Minor J.
ΑU
     Med. Sch., Univ. Michigan, Ann Arbor, MI, USA
CS
     Archives of Biochemistry and Biophysics (1976),
SO
     175(2), 524-33
     CODEN: ABBIA4; ISSN: 0003-9861
DT
     Journal
     English
LA
CC
     7-3 (Enzymes)
     Highly purified liver microsomal cytochrome P 450 (I) catalyzes the
AB
     hydroperoxide-dependent hydroxylation of a variety of substrates in the
     absence of NADPH, NADPH-I reductase, and mol. O. The addn. of
     phosphatidylcholine is necessary for maximal activity. The absence of
     flavoproteins and cytochrome b5 from the I prepns. rules out the
     involvement of other known microsomal electron carriers. The Fe+ form of
     I is not involved in peroxide-dependent hydroxylation reactions, as
     indicated by the lack of inhibition by CO. With cumene hydroperoxide (II)
     present, a variety of substrates is attacked, including N-methylaniline,
     N, N'-dimethylaniline, cyclohexane, benzphetamine, and aminopyrine. With
     benzphetamine as the substrate, II may be replaced by other peroxides,
     including H2O3, or by peracids or Na chlorite. A study of the
     stoichiometry indicated that equimolar amts. of N-methylaniline,
     formaldehyde, and cumyl alc. (.alpha.,.alpha.-dimethylbenzyl alc.) are
     formed in the reaction of N,N-dimethylaniline with II. Since H218O is
     incorporated only slightly into cyclohexanol in the reaction of
     cyclohexane with II, it appears that the O atom in cyclohexanol is derived
     primarily from the peroxide. The data obtained are in accord with a
     peroxidase-like mechanism for the action of I.
     cytochrome P450 substrate hydroxylation; cumene hydroperoxide cytochrome
ST
     P450
IT
     Hydroxylation
        (by cytochrome P 450, hydroperoxides in) .
TΤ
     Hydroperoxides
     RL: BIOL (Biological study)
        (in cytochrome P450 substrate hydroxylation)
                                     110-68-9 110-82-7, biological studies
ΙT
     58-15-1 100-61-8
                          103-67-3
     121-69-7
               156-08-1
                          935-67-1
     RL: BIOL (Biological study)
        (cytochrome P 450 action on)
     9035-51-2
ΙT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (hydroxylation by, hydroperoxides in)
TT
     80-15-9
     RL: BIOL (Biological study)
        (in cytochrome P 450 substrate hydroxylation)
=> d his
     (FILE 'HCAPLUS' ENTERED AT 16:37:35 ON 26 APR 2003)
                DEL HIS
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FILE 'REGISTRY' ENTERED AT 16:37:39 ON 26 APR 2003

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L1
              3 S 52-90-4 OR 921-01-7 OR 3374-22-9
                E N-ACETYL-L-CYSTEINE/CN
L2
              1 S E3
                E PANTOTHENIC ACID/CN
L3
              1 S E3
                E CHOLINE/CN
              1 S E3
T,4
                E THYMIDINE/CN
L5
              3 S E3
L6
             47 S C10H14N2O4/MF AND OC4/ES AND NCNC3/ES
L7
            154 S (3H OR H3) (L) THYMIDINE
1.8
              6 S L7 AND 2/NR
L9
              4 S L8 NOT P/ELS
L10
              7 S L6 AND (T OR D)/ELS
L11
              1 S 172294-32-5
L12
              5 S L9, L11
                E CUMENE HYDROPEROXIDE/CN
L13
              1 S E3
                E CHLORIDE, ION/CN
                E CHLORINE, ION/CN
L14
              2 S E9, E20
L15
             32 S CHLORINE AND ION AND 1/ATC AND 1/ELC.SUB
L16
             22 S L15 NOT ISOTOPE
L17
             20 S L16 NOT (37CL1 OR 35CL1)
                E PHOSPHATE, ION/CN
                E PHOSPHATE/CN
              1 S E3
L18
                E CALCIUM, ION/CN
L19
              2 S E10, E23
L20
             31 S CALCIUM AND ION AND 1/ATC AND 1/ELC.SUB
             22 S L20 NOT ISOTOPE
L21
                E MAGNESIUM, ION/CN
              2 S E4, E38
L22
L23
             29 S MAGNESIUM AND ION AND 1/ATC AND 1/ELC.SUB
L24
             15 S L23 NOT ISOTOPE
                E POTASSIUM, ION/CN
L25
              1 S E6
L26
             23 S POTASSIUM AND ION AND 1/ATC AND 1/ELC.SUB
L27
             20 S L26 NOT ISOTOPE
                E SODIUM, ION/CN
L28
              1 S E4
L29
             16 S SODIUM AND ION AND 1/ATC AND 1/ELC.SUB
L30
             13 S L29 NOT ISOTOPE
                E IRON, ION/CN
L31
              1 S E6
L32
             28 S IRON AND ION AND 1/ATC AND 1/ELC.SUB NOT ISOTOPE
L33
              6 S (CHLORINE OR CALCIUM OR MAGNESIUM OR POTASSIUM OR SODIUM OR I
L34
              1 S PHOSPHORIC ACID/CN
L35
              1 S WATER/CN
L36
              9 S C5H9NO3S/MF AND CYSTEINE AND ACETYL
L37
              3 S L36 NOT (14C# OR D/ELS OR 35S OR ESTER OR PROPANOIC)
     FILE 'HCAPLUS' ENTERED AT 16:54:20 ON 26 APR 2003
L38
           4758 S L2
           4777 S L37
L39
L40
           2346 S N ACETYL (1W) CYSTEINE
L41
           4056 S N ACETYLCYSTEINE
L42
             19 S L() (ACETYLCYSTEINE OR ACETYL CYSTEINE)
L43
           5363 S ACETYLCYSTEINE OR ACETYL CYSTEINE
           7201 S L38-L43
L44
           3439 S L3
L45
L46
          10269 S L4
          50295 S PANTOTHENIC ACID OR CHOLINE
L47
```

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L48
            224 S VITAMIN B5
L49
            314 S VITAMIN B3
L50
           3787 S PANTOTHENATE
          53009 S L46-L50
L51
           5128 S L14,L17,L18,L19,L21,L22,L24,L27,L25,L28,L39,L31,L32,L33,L34 A
L52
L53
          23705 S (CHLORIDE OR PHOSPHATE OR CALCIUM OR MAGNESIUM OR POTASSIUM O
L54
          23854 S L52, L53
           5456 S L13
L55
L56
           7373 S (CUMENYL OR CUMYL OR CUMEN#)()(HYDROPEROXIDE OR HYDRO PEROXID
L57
             68 S PERCUMYL H
            422 S PERCUMYL
L58
L59
            567 S ISOPROPYLBENZENE HYDROPEROXIDE
L60
             73 S CUMENEHYDROPEROXIDE OR CUMYLHYDROPEROXIDE OR CUMENYLHYDROPERO
L61
             10 S KAYACUMEN#
L62
              4 S L54 AND L55-L61
L63
              2 S L62 NOT (IDB OR HELMINTH)/TI
L64
             71 S L54 AND L44
L65
              1 S L64 AND L55-L61
              3 S L64 AND LYMPHOCYT?
L66
              2 S L66 NOT AZT
L67
L68
              3 S L63, L67
            360 S L1 AND L51
L69
            889 S CYSTEINE AND L51
L70
L71
              4 S L69, L70 AND L55-L61
L72
              2 S L71 AND L54
              3 S L68, L72
L73
L74
              2 S L71 NOT L73
L75
          21435 S (H3 OR 3H) (S) THYMIDIN?
            621 S L12
L76
           7249 S L5
L77
L78
           120 S L75-L77 AND L51
             48 S L78 AND L54
L79
L80
              0 S L78 AND L55-L61
L81
              1 S L79 AND LYMPHOCYT?
L82
             0 S L79 AND L44
L83
             16 S L79 AND L69, L70
L84
            ·13 S L83 AND (CULTUR? OR MEDIUM OR MEDIA)
                SEL DN AN 1 5-9
              6 S E1-E18
L85
L86
              9 S L68, L85
L87
             64 S L54 AND FREE (1A) SERUM
L88
             14 S L87 AND (L1 OR CYSTEINE)
L89
             17 S L86, L88
                SEL DN AN 3 8 17
L90
             14 S L89 NOT E19-E27
                SEL DN AN 2
L91
              1 S E28-E30
L92
             14 S L90, L91
                E CRAWFORD J/AU
            112 S E3, E11-E13
L93
             5 S E117
L94
L95
             41 S E107
L96
              2 S L93-L95 AND L44,L51
              2 S L93-L95 AND L55-L61
1.97
L98
              0 S L93-L95 AND L75-L77
L99
             15 S L92, L96, L97
     FILE 'HCAPLUS' ENTERED AT 17:20:44 ON 26 APR 2003
L100
             15 S L99 AND L38-L99
     FILE 'WPIX' ENTERED AT 17:21:43 ON 26 APR 2003
                E US20020068270/PN
L101
              1 S E3
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L102
           4556 S L47/BIX
            323 S L48/BIX OR L49/BIX OR L59/BIX
L103
           1352 S (R00467 OR R00231)/DCN OR (0467 OR 0231)/DRN
L104
           5335 S L102-L104
L105
           1739 S L56/BIX OR L57/BIX OR L58/BIX OR L59/BIX OR L60/BIX OR L61/BI
L106
L107
           1792 S R00474/DCN OR 0474/DRN
L108
            122 S L105 AND L106, L107
                E R04369+ALL/DCN
                E R11179+ALL/DCN
                E R06646+ALL/DCN
L109
           1610 S E1
                E R06671+ALL/DCN
            840 S E1
L110
                E R060311+ALL/DCN
                E R06031+ALL/DCN
L111
            816 S E1
                E R07107+ALL/DCN
L112
           1197 S E1
                E R06645+ALL/DCN
           1231 S E1
L113
                E R06108+ALL/DCN
L114
           1016 S E1
                E R04811+ALL/DCN
            941 S E1
L115
                E R04810+ALL/DCN
            830 S E1
1.116
L117
              2 S L108 AND L109-L116
              1 S L108 AND R04369/DCN
L118
            700 S L40/BIX OR L41/BIX OR L42/BIX OR L43/BIX
L119
              0 S L119 AND L108
L120
              1 S CYSTEINE/BIX AND L108
L121
L122
              2 S L117, L118, L121
              2 S C12N005/IC, ICM, ICS AND L108
L123
              0 S G01N003/IC, ICM, ICS AND L108
L124
              4 S C12Q001/IC, ICM, ICS AND L108
L125
L126
              4 S L122, L123, L125
L127
              2 S L126 AND CULTURE MEDIUM/TI
     FILE 'WPIX' ENTERED AT 17:33:14 ON 26 APR 2003
     FILE 'DPCI' ENTERED AT 17:33:26 ON 26 APR 2003
                E WO9810092/PN
L128
              1 S E3
                E WO9748821/PN
L129
              1 S E3
     FILE 'DPCI' ENTERED AT 17:34:10 ON 26 APR 2003
     FILE 'WPIX' ENTERED AT 17:36:54 ON 26 APR 2003
              8 S (US5985665 OR US4927762 OR US5171885 OR US5290571 OR US532669
L130
L131
              1 S US5985665/PN
L132
              7 S L130, L131 NOT L127
     FILE 'WPIX' ENTERED AT 17:38:01 ON 26 APR 2003
     FILE 'MEDLINE' ENTERED AT 17:38:31 ON 26 APR 2003
                E ARCH BIOCHEM BIOPHYS/JT
              1 S E3 AND 1976/PY AND NORDBLOM?/AU AND (175 AND 524)/SO
L133
                E ANAL BIOCHEM/JT
L134
              1 S E3 AND 1992/PY AND JIANG?/AU AND (202 AND 384)/SO
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FILE 'MEDLINE' ENTERED AT 17:39:43 ON 26 APR 2003

FILE 'HCAPLUS' ENTERED AT 17:39:56 ON 26 APR 2003 E ARCH BIOCHEM BIOPHYS/JT

1 S E3 AND NORDBLOM?/AU AND 1976/PY AND (175 AND 524)/SO

FILE 'HCAPLUS' ENTERED AT 17:40:23 ON 26 APR 2003